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(54) **Products comprising a sheet and a lipid and aqueous phase**

(57) This invention concerns products for cleansing and other applications that comprise a sheet of absorbent material, in particular of non-woven material, such as a wipe, to which are applied a lipid and an aqueous phase. The invention further concerns the manufacture and use of such products.

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Description

Field of the Invention

[0001] This invention concerns products for cleansing and other applications, which products comprise a sheet of absorbent material, in particular of a non-woven material, such as a wipe, to which a lipid and aqueous phase have been applied. The invention further concerns the manufacture and use of such products.

Background of the Invention

[0002] For a long time now, wipe products have been used as articles useful in cleansing in a wide range of applications while over the last two decades so-called wet wipes have become established as products suited for many personal cleansing applications. These products are typically manufactured by impregnating sheets made of non-woven fabric with a suitable liquid frequently referred to as lotion.

[0003] Recent innovations in the wipes area are based on improvements in the fabric, impregnate as well as product presentation. Initially, wet wipe products were made of traditional non-woven material based on paper making technology (pulp based products). These products were well accepted but deficient in softness of the fabric material. The introduction of the 'spunlace' non-woven technology offered a product which was superior in terms of the softness. An innovation step of another kind was the introduction of 'Pop-up' technology which offered a product superiority as regards the dispensing of individual wipes.

[0004] In addition to the above lotions have been developed which offer skincare benefits in addition to the basic cleansing properties of the wipe. One approach was the introduction of lotions that were based on oil-in-water emulsions which deliver superior mildness, moisturisation, protection and skin smoothness when compared to simple aqueous cleansing formulations. Other wet wipe products contain active skincare ingredients incorporated into simple aqueous formulations, which delivered soothing benefits, e.g. chamomile. Current wet wipe executions are simply impregnated with either aqueous lotions or with oil in water emulsions. Also the impregnation with oily formulations has been described.

[0005] However, these attempts have several limitations. Firstly, only a small part of the lotion (~15%) is released from the wipes during use. Thus a large quantity of the relatively expensive impregnate is not utilized to the benefit of the consumer since it never comes into contact with the skin and is therefore wasted when the product is discharged after use. This fact also prevents the use of expensive but more effective ingredients. Secondly, from a formulation point there is an apparent contradiction in the optimization of cleansing performance and skincare benefits in one single lotion since in-

gredients which are effective in cleansing may not be compatible with efficient skin care agents.

[0006] Another important factor in cleansing is the fact that a number of soils are water compatible and therefore more easily removed by water-based formulations, whereas others are lipid compatible and therefore adequately removed by lipid or oil based formulations. A complete and effective removal of soils therefore requires the presence in or on the wipe of as well water and oil-based components.

[0007] This is in particularly required in wipes for personal cleansing and in particular for wipes used for babies and infants. In the latter instance wet wipes are used for cleansing the perianal region when changing diapers. Inadequate cleaning not only results in personal discomfort but also gives rise to diaper rash and other infection related phenomena. It has been shown that the most effective way of preventing diaper rash is to cleanse the skin thoroughly and to remove the microorganisms that have been identified as causative. The source of these microorganisms is often the fecal deposits that can remain on a baby's skin while wearing the diaper. Because fecal deposits consist of both water-soluble and oil-soluble matter, however, complete removal of fecal deposits from the diaper area requires both water-based and oil-based cleansing agents.

[0008] Thus, it is an object of this invention to provide a mechanism for cleansing babies' skin in order both to remove waste deposits and to reduce the number of microorganisms available to cause infection.

[0009] WO 96/14835 discloses dry tissues to which a water-in-lipid emulsion has been applied and WO 99/25923 concerns a process and an apparatus for selectively coating a wipe with a water-in-lipid emulsion. WO 99/01536 discloses wipes wherein the carrier comprises two regions of different basis weight being applied with an emulsion comprised of a solidified lipid phase, a polar phase dispersed therein and an emulsifier. Other prior art in this field is WO 95/35411, WO 95/35412, WO 95/16824, WO 97/30216, DE 33 09 530 and the publication of R.E. Mathis in Nonwovens World 1999, pp. 59-65.

[0010] It is an object of this invention to offer a cleansing article and in particular a wet wipe product that allows to independently optimize the cleansing and skincare attributes of the product and at the same time improves the delivery of skincare actives onto the skin during use.

[0011] It is a further object of this invention to provide products that have an improved release of the active ingredient(s) onto the skin during use. It is still a further object of the present invention to provide a product for use as a cleansing tool that effectively and completely removes oil and water compatible soils.

[0012] Another object of this invention is to provide products for cleansing and other applications that allow convenient and quick application, are easy to carry, as well as an easier and more evenly distribution of the

product. They moreover should be easy to apply on babies and children.

[0013] These objects are attained by the products according to the present invention which comprise a sheet that contains an aqueous and a lipid phase. Whereas traditional wet wipes have been based on impregnation of a fabric with one phase, the products of this invention concern the use of two distinctly different impregnate phases (both in terms of physical appearance and location on the wipe) applied onto a single sheet. This approach allows optimizing the cleansing performance of the product independently from skincare properties.

Summary of the Invention

[0014] This invention relates to products that comprise a porous or absorbent sheet whereto a lipid and an aqueous phase have been applied. Preferably, the lipid phase is solid or semi-solid at ambient temperature and preferably is present at the surface or at the surface portion of one or both sides of the sheet.

In particular said sheet is made of a non-woven material, more in particular a non-woven material made by the spunlace or preferably the hydro-entanglement procedure.

[0015] In a further aspect there is provided a method of manufacturing a product as described herein, said method comprising applying to the sheet a lipid phase and an aqueous phase, either subsequently or simultaneously. In a preferred method of manufacturing, said sheet is first coated with a lipid phase and subsequently sprayed or impregnated with an aqueous phase.

[0016] In still a further aspect there is provided the use of a product as described herein as a cleansing tool, in particular in personal care applications.

[0017] In another aspect the invention concerns the use of a product as described herein as an applicator of active substances.

[0018] In still another aspect the invention provides the use of a product as described herein as a combined cleanser and applicator of active substances.

Detailed Description of the Invention

[0019] The absorbent or porous sheet can take the form of a tissue, a wipe, towel, towelette, and the like.

[0020] Materials of which the sheet is made may be mono or multi-layered, woven or non-woven. They can be made of one or of several materials. Particularly preferred are non-woven materials that have a web structure of fibrous or filamentous nature, in which the fibres or filaments are distributed randomly or with a certain degree of orientation, the former being obtainable by air-laying or certain wet-laying processes, the latter in other wet-laying or in carding processes. The fibres or filaments can be natural, for example wood pulp, wool cotton, linen and the like, or synthetic, for example polyvinyls, polyesters, polyolefins, polyamides and the like.

[0021] Typically they have a weight per square meter in the range of 10 to 80 g/m², in particular of 20 to 70 g/m². Particular materials are of the non-woven type. Based on the raw material that has been used, two different types of products can be distinguished.

[0022] A first type of carriers is paper based. The raw materials for these carriers are made almost exclusively of cellulose-based fibres or filaments from plant cellular sources (pulp). These can be available from fresh wood-shavings or from recycled material (recycled paper). In a number of wipe applications, such as baby wipes, wipes for cleansing, wet paper towels and the like, high wet strength or firmness of the non-woven web is a desirable attribute. This can be achieved by the addition of binding materials. Examples of such materials are the so-called wet strength resins. In some cases additives are added in order to increase the softness of the end product.

[0023] In a second type use the web is made mainly of staple, e.g. based on cotton, wool, linen and the like.

[0024] Commercial products are made of cellulose fibres, synthetic fibres or mixtures of both. Polyester and polypropylene are known as suitable polymers for the preparation of synthetic fibres. Also in these products binders can be used to increase the firmness of the non-woven fabric.

[0025] Webs of increased strength can be obtained by using the so-called spunlace or hydro-entanglement technique. In this technique the individual fibres are twisted together so that an acceptable strength or firmness is obtained without using binding materials. The advantage of the latter technique is the excellent softness of the non-woven material.

[0026] Non-woven materials that are made of a mixture of pulp and staple are also known. Such materials are available with binding materials, in particular those mentioned above, or without binding materials. In the latter instance the non-woven is preferably made by the spunlace or hydro-entanglement procedure.

[0027] In a preferred embodiment of the present invention, the carrier material is made of cellulose pulp with a small amount of binding material. The amount of binder in the carrier material is in the range of 5 to 20 % (w/w).

[0028] In a particularly preferred embodiment the non-woven fabric is prepared by the water entanglement procedure and does not contain binding material.

[0029] The absorbing ability of the carrier material is of particular interest with regard to the applications envisaged by the present invention. During production the impregnating solution should be taken up quickly by the carrier. In certain embodiments of this invention the wipes will be packed in a stack of a plurality of wipes. In this instance the absorbing ability of the non-woven fabric should be such that a chromatographic effect (sinking down of the lotion) in the stack is avoided during storage. On the other hand it should be guaranteed that during the usage of the wipe the impregnating solution is de-

livered evenly to the skin and the active ingredients are released quantitatively.

[0030] The absorbing capacity of the carrier material is determined essentially by three different parameters: the surface weight of the carrier material, the nature of the raw material used in the manufacture and the manufacturing process used.

[0031] For the applications according to the invention the carrier materials typically have a surface weight from 10 g/m² to 80 g/m², preferably from 30 to 70 g/m² and more preferably from 40 to 60 g/m². The selection of the raw material of which the non-woven carrier is made depends on the manufacturing procedure. Typically in the manufacture of non-woven carriers by the hydro-entanglement process, use is made of mixtures of cellulose fibres and synthetic fibres. The relative quantity of synthetic fibres in the non-woven fabric is from 0 to 100 % and preferably is between 10 and 70 %, more preferably in the range of 30 to 50 % (all percentages being w/w).

[0032] According to this invention the sheet material is contacted with a lipid and an aqueous phase.

[0033] Either of the phases may be liquid or semi-solid. Preferably, one phase is solid or semi-solid, e.g. a cream, while the other is liquid.

[0034] The phases may be applied to the whole sheet, i.e. continuously, or to parts of the sheet, i.e. discontinuously. One phase may be applied continuously while the other is applied discontinuously. They can be applied at the surface or in the internal of the sheet. If applied at the surface, one or both phases can be present at one side or at both sides of the sheet, or one phase may be present at one side while the other phase is present at the other side of the sheet.

[0035] In the instance where a phase or both phases are applied discontinuously, they are present at certain areas, in particular at one or more areas of the sheet. In that instance, the phase or phases may be present as one or more forms or shapes. For example they can be present as dots or spots, lines or stripes, as geometrical figures such as squares, rectangles, circles and the like, as symbols such as letters, text, logos, figures and the like, or as trademark signs, or any other such forms, or a combination thereof. The forms or shapes may be present over the entirety of the sheet or grouped in one or more areas, for example in a corner or in the center area.

[0036] In a particular embodiment, one phase is applied on one or on both sides of the sheet in the form of stripes, dots or other forms covering the entire surface or only a part of the surface of the sheet. The aqueous phase is applied to the sheet either on the entire surface of the fabric or on certain areas.

[0037] This may be done in a second step preferably after the application of the lipid phase or simultaneously in a one step operation.

[0038] In a preferred embodiment, both phases are applied subsequently to the sheet, more preferably first the lipid phase and subsequently the aqueous phase.

The lipid phase

[0039] A key feature of this invention is that the lipid phase applied to the sheet is formulated such that it is insoluble or at least essentially insoluble in the aqueous phase, although in some embodiments the two phases may be mixible or soluble into each other to a limited extend.

[0040] In a particular embodiment, the lipid phase is hydrophobic and is composed of materials that are generally insoluble in water such as oils or fats, or waxes.

[0041] The lipid phase preferably is solid or semi-solid at ambient temperature. It can take the form of a solid that is amorphous, semi-crystalline or crystalline, or it can take the form of a cream or waxy composition.

[0042] In particular the lipid phase has a melting point or a melting range above room temperature, in particular above 25 °C, for example in the range of 25 to 100°C, in particular in the range of 30 to 75°C, more in particular of 30 to 45°C, preferably in the range of 32 and 40°C. More preferably the melting temperature or melting range is above human body temperature. Most preferably the melting temperature or melting range approximates or is equal to human body temperature.

[0043] As used herein the term 'melting range' refers to a temperature range that starts from the temperature at which a substance or composition loses its solid constituency up to the temperature where it becomes completely liquid. A melting range is considered to be within a defined temperature range when it overlaps with that defined temperature range, or should be considered to be above a specified temperature when the range is above said temperature.

[0044] As used herein 'ambient temperature' refers to a temperature that is in the range of about 20 to about 25 °C.

[0045] Particularly preferred are the compositions of the lipid phase which are solid at room temperature and which have a penetration value of 0.2 - 4 mm (measured with: Petrotester PNR 10, Mikrokonus, 5 sec., temp 20 °C).

[0046] The water content of the lipid phase is low, in particular less than 10 %, preferably less than 6 %, more preferably less than 3 %. In a particular embodiment the lipid phase is water free, and will be such that it is not decomposed by the aqueous phase. As used herein, 'water free' means that the phase is composed of materials of low water content to which no water has been added.

[0047] The lipid phase may comprise one or more components selected from oils or fats, or waxes. It may further contain other components such as emollients. As used herein oils or fats refer to the same type of materials, oils being liquid at ambient temperature and fats being solid or semi solid at ambient temperature.

[0048] The lipid phase may also comprise mixtures of waxes and fats and/or oils.

[0049] In a preferred embodiment, the lipid phase is

a wax based composition, wherein the term 'wax' is as specified hereinafter.

Oils and fats

[0050] The lipid phase may contain oils, fats or mixtures of fats with oils and/or with oily components. The resulting mixture of which the lipid phase is composed should preferably selected such way that the melting point or melting range of the lipid phase is as mentioned above, in particular is above ambient temperature, more in particular is in the range of 32 °C to 40 °C.

[0051] Oils or fats which can be used in the lipid phase comprise natural oils or fats, or natural oil or fat derivatives, in particular of vegetable origin. Examples are almond oil, soybean oil, sunflower oil, safflower oil, corn oil, kernel oil, canola oil, borage oil, evening primrose oil, grapeseed oil, wheat germ oil, avocado oil, jojoba oil, sesame oil, walnut oil, linseed oil, palm oil, olive oil, macadamia oil, castor oil, rapeseed oil, peanut oil, coconut oil, and turnip seed oil, and the hardened derivatives thereof. The latter are obtained by hydrogenation of fats or oils. Preferred are hardened oils or fats from vegetal origin, e.g. hardened castor oil, peanut oil, soya oil, turnip seed oil, cotton seed oil, sunflower oil, palm oil, kernel oil, linseed oil, almond oil, corn oil, olive oil, sesame oil, cocoa butter, shea butter and coconut oil.

[0052] Said hardened fats or oils have the additional advantage of increasing the constituency of the lipid phase compositions.

[0053] The lipid phase may further comprise fatty components isolated from these natural oils, i.e. pure triglycerides or mixtures thereof, or the latter components having been prepared chemically. These so-called triglycerides (or triacyl glycerines) are esters of glycerines with fatty acids or fatty acid mixtures, for example so called technical mixtures obtained by hydrolysis from fractions of oils or fats, or by fractioning fatty acid mixtures after hydrolysis. The triglycerides may also be obtained chemically by synthesis.

[0054] The fatty acids in said triglycerides may be saturated or unsaturated, straight or branch chained, substituted or unsubstituted. Preferred triglycerides are those glycerines esters derived from fatty acids, either saturated or unsaturated, having from 10 to 60, in particular from 12 to 36, more particularly from 12 to 24, preferably from 16 to 20 carbon atoms. Preferred such fatty acids are, for example, palmitic, palmitic, oleic, lauric, myristic, stearic, hydroxystearic, behenic acid, or mixtures thereof. Within this group the triglycerides derived from saturated fatty acids are of particular interest.

[0055] Of particular interest are glyceryl tristearate, also referred to as stearin, glycerine tribehenate, glycerine tripalmitate, glycerine trilaurate, glycerine trioleate, glycerine trimyristate.

[0056] The lipid phase may also contain mono- or diglycerides, optionally in a mixture with the fats and oils mentioned herein, in particular with triglycerides. The

mono- or diglycerides for use in the lipid phase are derived from saturated or unsaturated, linear or branch chained, substituted or unsubstituted fatty acids or fatty acid mixtures. Also in this instance the melting point or melting range of the lipid phase preferably is as mentioned above, in particular is above ambient temperature, more in particular is in the range of 32 °C to 40 °C. Particular mono- or diglycerides are mono- or di-C₁₂₋₂₄ fatty acid glycerides, specifically mono- or di-C₁₆₋₂₀ fatty acid glycerides, for example glyceryl monostearate, glyceryl distearate. Mixtures of mono-, di- and, optionally, triglycerides can be derived from fractions of fatty acids. An example of such mixture for use as a component of the lipid phase is a mixture of C₁₂₋₁₈ mono-, di- and triglycerides.

[0057] In a preferred embodiment according to the present invention the lipid phase contains one or more fatty acid glycerides selected from the mono-, di- or triesters from glycerine, or a mixture thereof. The glycerides can be present in various amounts, it is typically present in an amount of up to 60% or in certain embodiments up to 70 %, or up to 80 % (w/w).

[0058] In other embodiments, in particular those containing dialkyl(ene)ethers or - carbonates, dicarboxylic acids or hydroxy fatty alcohols, the amount of said fatty ester glycerides will be up to 50 % and more preferably up to 40 % (w/w), relative to the total quantity of the lipid phase.

[0059] In a particular aspect of this invention there provided products as specified herein wherein the lipid phase consists essentially of one or more fatty acid glycerides selected from the mono-, di- or triesters from glycerine, or a mixture thereof. The glyceride can be present in various amounts, e.g. the amounts mentioned hereinabove or hereinafter.

[0060] Mixed esters as well as mixtures of mono-, di- and triglycerides are of particular interest because of their low propensity to crystallize and their capacity to improve the constituency of the formulation making up the lipid phase.

[0061] The lipid phase may also comprise alkyl esters of fatty acids, wherein the alkyl group has from 1 to 30 carbon atoms, preferably from 12 to 24 carbon atoms. The fatty acids in said alkyl esters in particular are C₁₂₋₃₀ fatty acids, more in particular C₁₂₋₂₀ fatty acids. The alkyl groups in said esters preferably are derived from fatty alcohols as well as of mixtures thereof, which, for example, are obtained by high pressure hydrogenation of technical mixtures of the methyl esters derived from fats or oils.

[0062] Preferred are the alkyl esters of C₁₆₋₂₄ fatty acids, more preferably from C₁₆₋₁₈ fatty acids, and C₁₋₃₀ fatty alcohols, preferably C₈₋₂₄ fatty alcohols, more preferably C₁₂₋₂₀ fatty alcohols.

[0063] Of particular interest in this regard are, e.g. stearyl stearate, palmityl stearate, stearyl behenate, cetyl stearate, cetyl behenate, cetyl palmitate, cetearyl behenate, behenyl behenate, stearyl heptanoate, stearyl

octanoate, myristyl myristate, myristyl isostearate, myristyl oleate, cetyl isostearate, cetyl oleate, stearyl isostearate, stearyl oleate, isostearyl myristate, isostearyl palmitate, isostearyl stearate, isostearyl isostearate, isostearyl oleate, isostearyl behenate, isostearyl oleate, oleyl myristate, oleyl palmitate, oleyl stearate, oleyl isostearate, oleyl oleate, oleyl behenate, oleyl erucate, behenyl isostearate, behenyl oleate, erucyl isostearate.

[0064] Of further interest are esters of linear C_6 - C_{22} -fatty acids with branched alcohols, in particular 2-ethylhexanol, esters of branched C_6 - C_{22} -fatty acids with linear alcohols, esters of C_{18} - C_{38} -alkylhydroxycarbonic acids with linear or branched C_6 - C_{22} -fatty alcohols, esters of linear and/or branched fatty acids with polyalcohols (e.g. propylene glycol, dimerdiol or trimetriol) and/or Guerbet alcohols, as well as esters of C_6 - C_{22} -fatty alcohols and/or Guerbet alcohols with aromatic carbonic acids, in particular benzoic acid, esters of C_2 - C_{12} -dicarbonic acids with linear or branched C_1 - C_{22} -alcohols (e.g. dioctyl malate) or C_2 - C_{10} -polyols having 2 to 6 hydroxyl groups.

[0065] Preferred fats comprise the triglycerides, in particular those derived from fatty acids having from about 12 to about 24 carbon atoms, in particular those having from about 12 to about 20 carbon atoms, more in particular those having from about 16 to about 20 carbon atoms. These fatty acids may be unsaturated or, which is preferred, saturated.

[0066] Particularly preferred are glycerides derived from oleic, stearic, myristic or lauric acid, or from fatty acid mixtures derived from natural oils such as cocoacids. Examples of preferred fats are cocoglycerides, glyceryl stearate, glyceryl laurate, and the like.

[0067] Further preferred fats comprise hydrogenated natural oils such as hydrogenated castor oil, hydrogenated palm oil and the like.

[0068] The lipid phase may also comprise oily components, i.e. non water-mixible components that are liquid at 20 °C. These can be e.g. glycerides, hydrocarbons, silicon oils, ester oils and the like, as well as mixtures thereof. The total quantity of these oily components in the total composition of the lipid phase preferably will be such that the lipid phase is solid at room temperature, or that it has a melting point or range that is as specified hereinabove. The oily components will typically be present in quantities of less than 40 % (w/w), in particular less than 20 %, or further in particular 1-15 %, more in particular from 2-10 % (w/w) relative to the total quantity of the lipid phase.

[0069] The oily components can be any of the oils mentioned hereinabove as 'oils and fats', more in particular the mono-, di- and triglycerides mentioned hereinabove, that are liquid at 20 °C. The oily components can further be fatty acids and fatty alcohols, described hereinabove in the respectively sections, therein that are liquid at 20 °C.

[0070] Further oily components which can be used in the lipid phase comprise silicone oils, mineral and par-

affin oils and synthetic oils, either aliphatic or aromatic, as well as mixtures thereof. Examples of such oils are squalane, squalene, isohexadecane, isoeicosane, polydecene, and also members of the group of dialkylcyclohexanes.

[0071] The lipid phase may further contain silicone oils such as, for example cyclic silicones, dialkyl- or alkylarylsiloxanes, e.g., cyclomethicone, dimethyl polysiloxane and methylphenyl polysiloxane, as well as the alkoxyated and quaternized analogs thereof. Appropriate non-volatile silicon oils are e.g. polyalkylsiloxanes, polyalkylarylsiloxanes and polyethersiloxane-copolymers.

[0072] The total amount of fats or oils, or of mixtures of fats and oils and/or oily components in the lipid phase in particular is at least 50 %, preferably at least 70 %, more preferably at least 90 %, w/w of the total amount of components making up the lipid phase.

[0073] In a particular aspect of this invention there are provided products as specified herein wherein the lipid phase essentially consists of fats or oils, or of mixtures of fats and oils and/or oily components, in particular those specified in this specification. The fats, oils and oily components can be present in various amounts, e.g. the amounts mentioned hereinabove or hereinafter.

Waxes

[0074] The lipid phase may comprise waxes. As used herein, the term 'wax' refers to oil soluble materials that have a waxy constituency and have a melting point or range of above ambient temperature, in particular above 25 °C. Waxes are materials that have a solid to semi-solid (creamy) consistency, crystalline or not, being of relative low viscosity a little above their liquefying point. Waxes can be composed of one or more components, synthetic as well as natural, and can in principle be composed of or comprise any oil soluble material having a waxy constituency, including mixtures thereof.

[0075] Waxes which can be used may be synthetic or natural waxes, as well as other oil soluble materials that have a waxy consistency. Waxes also encompass materials such as oils or fats of natural or synthetic origin, and waxy components such as higher alkanols (in particular fatty alcohols), higher alkanediols (in particular hydroxy fatty alcohols) carboxylic acids (in particular fatty acids), dialkyl(ene)ethers, dialkyl(ene) carbonates, dicarboxylic acids and the like components.

[0076] Natural waxes comprise waxes from vegetal origin, such as purcelline, shea butter, cocoa butter, Japan wax, esparto gras wax, cork wax, Guaruma wax, rice shoot wax, Ouricury wax, montan wax, sunflower wax, ceresine wax, sugar cane wax, carnauba wax, candellilla wax, lanolin, fruit-derived waxes, such as orange wax, lemon wax, grapefruit wax and bayberry wax, and the like and of animal origin such as beeswax, woolwax, spermateci and bear fat, shellac wax, and the like. Natural waxes further comprise mineral waxes such as cer-

esine and ozokerite waxes. Synthetic waxes comprise petroleum-based waxes such as paraffin, vaseline, petrolatum, micro wax. Further synthetic waxes are polyalkylene and polyethyleneglycol waxes, e.g. polyethylene wax; waxes based on chlorinated naphthalenes such as 'Halowax', synthetic hydrocarbon waxes, and the like, including mixtures thereof. Further waxes are chemically modified waxes, in particular hardened or hydrogenated waxes such as, for example, Montan-ester waxes, Sasol waxes and hydrogenated jojoba waxes.

[0077] Preferred among these natural waxes are waxes from vegetal origin.

[0078] Other wax components can be certain fats (including mono-, di- and triglycerides and fatty acid alkylesters), fatty alcohols, fatty acids, including substituted fatty acids (in particular hydroxy substituted fatty acids, for example, 12-hydroxystearic acid), dialkyl(ene) ethers, dialkyl(ene) carbonates, dicarboxylic acids (in particular the C₁₆-C₄₀-dialkylesters of dicarboxylic acids, e.g. the C₁₆-C₄₀-alkyl stearates, C₁₈-C₃₈-alkylhydroxystearyl stearates or C₂₀-C₄₀-alkyl erucates) and hydroxy fatty alcohols that comply with the definition of 'wax' as outlined herein. Any of these components may contain homologous components that are liquid, as long as the total composition making up the lipid phase has a waxy constituency. For example, waxy fats may contain oils, waxy fatty alcohols may contain liquid fatty alcohols, etc., in such amount that the total composition has a waxy constituency and in particular has the melting point or range specified above.

[0079] Still further wax components are selected from the group of aromatic carbonic acids, tricarboxylic acids, or from the group of lactides of long-chained hydroxycarbonic acids. Myristyl lactate is particularly attractive for use on wipes for skin treatment, because of its binding capacity to the skin

[0080] Further wax components that can be used are C₃₀-C₅₀-alkyl bees wax; tri-C₁₆-C₄₀-alkyl citrates, e.g. tristearyl citrate, triisostearyl citrate, trilauryl citrate; ethyleneglycol di fatty acid esters, in particular the ethylene glycol di-C₁₂-C₃₀-fatty acid esters, e.g. ethylene glycol dipalmitate, ethyleneglycol distearate, ethyleneglycol di (12-hydroxystearate).

[0081] As further useful components there can be mentioned silicon waxes.

[0082] The lipid phase may also comprise mixtures of waxes and fats and/or oils.

[0083] The total amount of waxes in the lipid phase in particular is at least 50 %, preferably at least 70 %, more preferably at least 90 %, w/w of the total amount of components making up the lipid phase.

[0084] In a particular aspect of this invention there provided products as specified herein wherein the lipid phase essentially consists of one or more waxes selected from the waxes mentioned herein, including mixtures thereof. The waxes can be present in various amounts, e.g. the amounts mentioned hereinabove or hereinafter.

Fatty alcohols

[0085] The lipid phase may also comprise fatty alcohols. Fatty alcohols that can be used are, for example, C₁₂-C₅₀-fatty alcohols, in particular the C₁₂-C₂₄-fatty alcohols, that are derived from natural fats, oils or waxes such as, for example, myristylalcohol, 1-pentadecanol, cetylalcohol, 1-heptadecanol, stearylalcohol, 1-nonadecanol, arachidylalcohol, 1-heneicosanol, behenylalcohol, brassidylalcohol, lignocerylalcohol, cerylalcohol or myricylalcohol as well as Guerbet alcohols. Preferred for use in the present invention are saturated, straight or branch chained fatty alcohols. However also unsaturated, straight or branch chained alcohols can be used, optionally in a mixture with saturated alcohols. Preferably the alcohols will be selected such that the melting point of the mixture is as referred to hereinabove and more in particular is in the range of 32 to 40 °C.

[0086] Mixtures of fatty alcohols can evidently also be used, including fatty alcohol fractions obtained from the reduction of the corresponding fatty acid fractions derived from naturally occurring oils or fats such as, for example, almond oil, soybean oil, sunflower oil, safflower oil, corn oil, canola oil, borage oil, evening primrose oil, grapeseed oil, wheat germ oil, avocado oil, jojoba oil, sesame oil, walnut oil, linseed oil, palm oil, olive oil, castor oil, macadamia oil, rapeseed oil, peanut oil, coconut oil, and turnip seed oil.

[0087] Synthetic alcohols can also be used such as, for example, the linear fatty alcohols of an even number of carbon atoms resulting from the Ziegler-synthesis (Alfole®) or the partially branched alcohols resulting from the Oxo synthesis (Dobanole®).

[0088] A preferred embodiment according to the present invention is that wherein the lipid phase contains at least one fatty alcohol, more preferably at least one C₁₄-C₁₈-fatty alcohol. Also preferred is a lipid phase with at least one C₁₆/C₁₈-Guerbet alcohol.

[0089] The use of fatty alcohols advantageously results in the lipid phase having a drier, i.e. less greasy, skin feel, compared to components such as triglycerides.

[0090] The total amount of fatty alcohols in the lipid phase may vary, and depends on the desired properties of the lipid phase. In a number of instances it is desirable to have a relative higher quantity of fatty alcohols in the composition, in particular said alcohols will be present in an amount of 50 %, preferably at least 70 %, more preferably at least 90 %, (w/w) of the total amount of components making up the lipid phase. In other instances, relatively lower amounts are desired, the total amount of the fatty alcohols present in the lipid phase is in the range of 1 - 40 %, preferably of 1 - 30 % (w/w), more preferably of 1 - 20 % (w/w), still more preferably from 1 - 10 % (w/w).

[0091] In a particular aspect of this invention there provided products as specified herein wherein the lipid phase essentially consists of one or more fatty alcohols,

in particular those specified in this patent specification, including mixtures thereof. The fatty alcohols can be present in various amounts, e.g. the amounts mentioned hereinabove or hereinafter.

Fatty acids

[0092] The lipid phase may also contain C₁₄-C₄₀-fatty acids, including mixtures thereof. Of particular interest are the C₁₆-C₃₀-fatty acids. These comprise, for example, myristic-, pentadecanoic-, palmitic-, margaric-, stearic-, nonadecanoic-, arachic-, behenic-, lignoceric-, cerotic-, melissic-, erucaic-, elaeostearic-, oleic-, lonolenic-, lauric acid as well as substituted fatty acids, e.g. hydroxy-substituted fatty acids such as, for example, 12-hydroxystearic acid, and the amide or monoethanolamides of these fatty acids.

[0093] The total amount of the C₁₄-C₄₀-fatty acids present in the lipid phase, relative to the total weight amount of the lipid phase, is in the range of 1 - 30 % (w/w), preferably of 1 - 20 % (w/w), more preferably from 1 - 10 % (w/w).

[0094] In a particular aspect of this invention there are provided products as specified herein wherein the lipid phase essentially consists of one or more fatty acids, in particular those specified in this patent specification, including mixtures thereof. The fatty acids can be present in varying amounts, e.g. the amounts mentioned hereinabove or hereinafter.

Dialkyl(ene)ethers or -carbonates, dicarboxylic acids or hydroxy fatty alcohols

[0095] The lipid phase may also contain dialkyl(ene) ethers, dialkyl(ene) carbonates, dicarboxylic acids or hydroxy fatty alcohols, or mixtures thereof, which ethers, carbonates, acids or alcohols in particular those described hereinafter.

[0096] In a particular aspect of this invention there are provided products as specified herein wherein the lipid phase essentially consists of one or more dialkyl(ene) ethers or -carbonates, dicarboxylic acids or hydroxy fatty alcohols, including mixtures thereof. The dialkyl(ene) ethers or -carbonates, dicarboxylic acids or hydroxy fatty alcohols can be present in various amounts, e.g. the amounts mentioned hereinabove or hereinafter.

[0097] The addition of dialkyl(ene) ethers or -carbonates, dicarboxylic acids or hydroxy fatty alcohols, including mixtures thereof to the composition of the lipid phase allows to optimize the properties of the lipid phase, in particular its sensoric properties, i.e. the products as well as the skin after the products have been applied have a less greasier feel and also a less dry skin-feel, while having excellent skin caring properties.

Dialkyl(ene) ethers

[0098] The dialkyl(ene) ethers are symmetric or

asymmetric, straight or branch chained, saturated or unsaturated. Preferred are waxy, saturated C₁₆-C₃₀-dialkylethers, in particular C₁₆-C₂₄-dialkylethers. More preferred are C₁₆-C₂₀-dialkylethers, and particularly preferred are distearylethers and dibehenylethers. Dialkylethers of shorter chain length can also be used such as, for example, di-n-octylether, di-(2-ethylhexyl)-ether, laurylmethylether or octylbutylether, didodecylether, under the condition that the complete composition of the lipid phase has the desired melting point.

[0099] These ethers can be obtained from the appropriate fatty alcohols in the presence of an acid catalyst following art-known procedures. Typical examples are the products that are obtained by the etherification of capron alcohol, capryl alcohol, 2-ethylhexyl alcohol, caprin alcohol, lauryl alcohol, myristyl alcohol, cetyl alcohol, palmoleyl alcohol, stearyl alcohol, isostearyl alcohol, elaidyl alcohol, petroselinyl alcohol, linolyl alcohol, linolenyl alcohol, oleyl alcohol, rizinol alcohol, elaeostearyl alcohol, arachidyl alcohol, gadoleylalcohol, behenyl alcohol, erucyl alcohol and brassidyl alcohol, Guerbet alcohols, as well as mixtures thereof, which, for example, are obtained by high pressure hydrogenation of technical mixtures of the methyl esters derived from fats or oils.

[0100] Of particular interest are the dialkyl(ene) ethers that are solid at 25 °C.

Dialkyl(ene) carbonates

[0101] The dialkyl(ene) carbonates are symmetric or asymmetric, straight or branch chained, saturated or unsaturated. Preferred dialkyl(ene) carbonates are waxy, linear or branch chained, saturated or unsaturated C₁₄-C₃₀-dialkyl(ene) carbonates. More preferred are C₁₆-C₂₄-dialkyl carbonates and amongst these the saturated linear C₁₆-C₂₂-dialkyl carbonates. Particularly preferred is distearyl carbonate. Also liquid dialkyl(ene) carbonates, such as, for example, dihexyl-, dioctyl-, di-(2-ethylhexyl)- or dioleylcarbonate, can be used, under the condition that the complete composition has the desired melting point.

[0102] These dialkyl(ene) carbonates can be obtained by re-esterification of dimethyl- or diethylcarbonates with the corresponding hydroxy compounds following art-known procedures. Typical examples of dialkyl(ene) carbonates are re-esterification products of dimethyl- and/or diethylcarbonate with capron alcohol, capryl alcohol, 2-ethylhexyl alcohol, caprin alcohol, lauryl alcohol, myristyl alcohol, cetyl alcohol, palmoleyl alcohol, stearyl alcohol, isostearyl alcohol, elaidyl alcohol, petroselinyl alcohol, linolyl alcohol, linolenyl alcohol, oleyl alcohol, rizinol alcohol, elaeostearyl alcohol, arachidyl alcohol, gadoleyl alcohol, behenyl alcohol, erucyl alcohol and brassidyl alcohol, Guerbet alcohols, as well as technical mixtures thereof, that can be obtained by hydration of methyl esters derived from suitable oils or fats or oil or fat fractions.

[0103] Of particular interest are those dialkyl(ene) carbonates that are solid at 25 °C.

Dicarboxylic acids

[0104] Dicarboxylic acids that can be used are, for example, C₉-C₃₄-dicarbonic acids.

Hydroxy fatty alcohols

[0105] The hydroxy fatty alcohols for use in the said preferred or particularly preferred waxy compositions are saturated or unsaturated, straight chain or branched. Preferred are C₁₂-C₃₀-hydroxy fatty alcohols, at which the position of the hydroxy-substituent depends upon the synthesis route and the starting materials that have been used. The encompass for example 1,10-decanediol, 1,2-hexadecanediol, 12-

hydroxystearyl alcohol or hydroxy-Guerbet alcohols. Preferred are those hydroxy fatty alcohols that are solid at 25 °C, although liquid analogs can also be used, as long as the complete composition has the desired melting point. Particularly preferred is 12-hydroxystearyl alcohol.

[0106] The total amount of one or more of the dialkyl ethers, dialkyl carbonates, dicarbonic acids and the hydroxyalcohols present in the lipid phase, relative to the total weight amount of the lipid phase, is in the range of 1 - 30 % (w/w), preferably of 1 - 20 % (w/w) more preferably from 1 - 10 % (w/w).

Other components

[0107] The compositions of the lipid phase may contain further components, which may be of waxy nature or otherwise. The use of these further components allows to influence the sensorial properties as well as the stability of the compositions, in particular when after application to wipe material and more in particular when in contact with the aqueous phase. The other components may also be added to influence constituency, feel and appearance. These components will generally be insoluble or poorly soluble in water. Water soluble components can also be included, typically in combination with a solubilizing or emulsifying agent and some water.

Examples of further components are superfatting agents, thickeners, polymers, active ingredients, film forming agents, UV-filters, anti-oxidants, hydrotropic agents, preservatives, insect repellents, self-tanning agents, solubilizers, perfume oils, dyestuffs and the like.

[0108] Substances that can be used as **superfatting agents** are, for example, lanolin or lanolin derivatives such as lanolin alcohols, lanolin acids, polyethoxylated or acylated lanolin, or other lanolin derivatives; phospholipids such as lecithin or lecithin derivatives such as polyethoxylated or acylated lecithin or other lecithin derivatives; polyol fatty acid esters, monoglycerides and fatty acid alkanolamides.

[0109] Appropriate **thickeners** for example are of the Aerosil ®-type (hydrophile silica acids), polysaccharides, in particular xanthan-gum, guar-guar, agar-agar, alginate and tyloses, carboxymethyl cellulose and hydroxyethyl cellulose, additionally relatively high molecular weight polyethylene glycol mono- and -diesters of fatty acids, polyacrylate, (for example Carbopole® of Goodrich or Synthalene® of Sigma), polyacrylamides, polyvinylalcohol and polyvinylpyrrolidone, Surfactants such as, for example, ethoxylated fatty acid glycerides, ester of fatty acids with polyols such as, for example, pentaerythritol or trimethylolpropane, fatty alcohol ethoxylates with limited spread of homologs or alkyloligogluco-sides as well as electrolytes such as sodium chloride ammonium chloride.

[0110] Appropriate **cationic polymers** are for example cationic cellulose derivatives, e.g. quaternized hydroxyethyl cellulose (commercialized under the trade name Polymer JR 400® by Amerchol), cationic starches, copolymers of diallylammonium salts and acrylamides, quaternized vinylpyrrolidone/vinylimidazol-polymers (for example Luviquat® of BASF), condensation products of polyglycols and amines, quaternized collagen polypeptides, such as, for example, lauryldimonium hydroxypropyl hydrolyzed collagen (Lamequat® L/Grünau), quaternized wheat polypeptides, polyethylene imines, cationic silicone polymers, e.g., amidomethicone, copolymers of adipinic acid and dimethylamino-hydroxypropyldiethylenetriamine (Cartaretine® /Sandoz), copolymers of acryl acid with dimethyldiallylammonium-chloride (Merquat® 550/Chemviron), polyaminopolyamides, cationic chitine derivatives such as, for example, quaternized chitosans, optionally dispersed in mikrocristalline form, condensation products derived from dihalogenalkylenes, such as, for example dibromobutane with bis-dialkylamines, e.g. bis-dimethylamino-1,3-propane, cationic guar-gum, such as, for example, Jaguar® CBS, Jaguar® C-17, Jaguar® C-16 from Celanese, quaternized ammonium salt-polymers, e.g. Mirapol® A-15, Mirapol® AD-1, Mirapol® AZ-1 from Miranol.

[0111] **Anionic, zwitterionic, amphoteric and non-ionic polymers** that can be used are, for example, vinylacetate/crotonic acid-copolymers, vinylpyrrolidone/vinylacrylate-copolymers, vinylacetate/butylmaleate/ isobornylacrylate-copolymers, methylvinylether/maleic acid anhydride-copolymers and their esters, which are not cross-linked and with polyols linked polyacrylids which are cross-linked, acrylamidopropyl trimethylammonium chloride/ acrylate-copolymers, octylacrylamide/methylmethacrylate/tert.butylaminoethylmethacrylate/2-hydroxypropylmethacrylate-copolymers, polyvinylpyrrolidone, vinylpyrrolidone/vinylacetate-copolymers, vinylpyrrolidone/ dimethylaminoethylmethacrylate/vinyl caprolactam-terpolymers as well as optionally derivatized cellulose ethers and silicones.

[0112] As further **consistency agents** there can be used small amounts of alkalimetal or alkaline earth met-

al as well as aluminium salts of C₁₂-C₂₄-fatty acids or C₁₂-C₂₄-hydroxyfatty acids, preferred being calcium-, magnesium-, aluminium- and in particular zinc stearates.

[0113] The lipid phase may further contain suitable **anti-oxidants** such as, for example, sulfites, e.g. sodium sulfite, tocopherol or derivatives thereof, ascorbic acid or derivatives thereof, citric acid, propyl gallate, chitosan glycolate, cysteine, N-acetyl cysteine plus zinc sulfates, thiosulfates, e.g. sodium thiosulfates, polyphenols and the like.

[0114] The lipid phase may further contain powders or powdered ingredients or mixtures thereof such as talcum, Bolus alba, myristyl alcohol, cetyl alcohol, cetyl-stearyl alcohol, calcium or magnesium stearate, magnesium lauryl sulfate, starch or derivatives thereof e.g. distarch phosphate, aluminium starch octenylsuccinate, carboxymethyl starch, tapioca starch, dimethylimidazolidinone rice starch, sodium starch glycolate, potato starch, rice starch, corn starch, hydroxypropyl starch, hydroxyethyl starch and the like.

Preferred compositions

[0115] Preferred embodiments of the present invention are those wherein the lipid phase has the composition as described under I, II, or III hereinafter.

[0116] In a preferred embodiment, the composition of the lipid phase will in particular have a melting point or range of above 25 °C, preferably in the range of 30 to 45 °C, more preferably in the range of 32 to 40 °C.

[0117] The water content of the preferred compositions of the lipid phase is low, e.g. lower than 10 %, preferably lower than 6 %, more preferably lower than 3 %. In particular, the preferred compositions will be water free.

[0118] In a preferred embodiment I of the present invention, the lipid phase contains one or more fatty acid mono-, di- or triglycerides, or natural oils comprising mono-, di- or triglycerides as well as the hydrogenated derivatives of said natural oils. The fatty acids in said glycerides may be synthetic or derived from natural oils, including hydrogenated derivatives thereof. The fatty acids contain from 12 to 24, preferably from 16 to 20 carbon atoms.

[0119] A particular example of a hydrogenated derivative of a natural oil is hydrogenated castor oil.

Preferred embodiment I

[0120] In a particularly preferred embodiment I, the lipid phase comprises one or more mono-, di- or triglycerides, in particular a C₁₂₋₂₄ fatty acid mono-, di- or triglyceride, or more in particular a C₁₆₋₂₀ fatty acid mono-, di- or triglyceride. In still a further particularly preferred embodiment I, the lipid phase comprises one or more triglycerides, in particular a C₁₂₋₂₄ fatty acid triglyceride, or more in particular a C₁₆₋₂₀ fatty acid triglyceride. Par-

ticular examples of such triglycerides are glyceryl stearate, glyceryl oleate, glyceryl laurate, glyceryl myristate, cocoglycerides, hydrogenated palm glycerides.

[0121] The total amount of mono-, di- or triglyceride(s) in the lipid phase of the preferred embodiments I in particular is at least 50 %, preferably at least 70 %, more preferably at least 90 %, w/w of the total amount of components making up the lipid phase. More preferably, the total amount of triglyceride(s) in the lipid phase of the preferred embodiments I in particular is at least 50 %, preferably at least 70 %, more preferably at least 90 %, w/w of the total amount of components making up the lipid phase.

Preferred embodiment II

[0122] In a preferred embodiment II, the lipid phase contains C₁₂-C₅₀-fatty alcohols, in particular the C₁₂-C₂₄-fatty alcohols, that are derived from natural fats, oils or waxes such as, for example, myristyl alcohol, 1-pentadecanol, cetyl alcohol, 1-heptadecanol, stearyl alcohol, lauryl alcohol, oleyl alcohol, palmityl alcohol, cetearyl alcohol, 1-nonadecanol, arachidyl alcohol, 1-heneicosanol, behenyl alcohol, brassidyl alcohol, lignoceryl alcohol, ceryl alcohol or myricyl alcohol as well as Guerbet alcohols.

[0123] Of particular interest for use in the invention are C₁₄-C₁₈-fatty alcohols as well as C₁₆/C₁₈-Guerbet alcohols.

[0124] The total amount of one or more of the C₁₂-C₅₀-fatty alcohols present in the lipid phase, relative to the total weight amount of the lipid phase, is in the range of 1 - 30 % (w/w), preferably of 1 - 20 % (w/w) more preferably from 1 - 10 % (w/w).

Preferred embodiment III

[0125] In a preferred embodiment III the lipid phase is a waxy composition comprising: at least one oil or wax component selected from dialkyl(ene) ethers, dialkyl(ene) carbonates, dicarboxylic acids or hydroxy fatty alcohols or mixtures thereof.

[0126] In a particularly preferred embodiment III the lipid phase is a waxy composition comprising:

(a) at least one oil or wax component selected from dialkyl(ene) ethers, dialkyl(ene) carbonates, dicarboxylic acids or hydroxy fatty alcohols or a mixture thereof;

(b) an active ingredient.

[0127] Particular dialkyl(ene) ethers, dialkyl(ene) carbonates, dicarboxylic acids or hydroxy fatty alcohols for use in the lipid phase of preferred embodiment III are those mentioned hereinabove.

[0128] The said preferred or particularly preferred waxy composition preferably liquifies above 25 °C and/or has a water content of less than 10 %, preferably less

than 6 %, more preferably less than 3 %. In particular said preferred or further preferred waxy composition is water free, and will be such that it is not decomposed by the aqueous phase. As used herein, water free generally means that the phase is composed of materials of low water content to which no water has been added.

[0129] The lipid phase having the preferred composition III can contain the same further ingredients as those described in relation to the lipid phase, in particular further waxy lipid components or oils.

[0130] The lipid phase having the preferred composition III can also contain liquid dialkyl(ene) ethers, dialkyl(ene) carbonates, dicarbonic acids or hydroxy fatty alcohols, but preferably in such amounts that the melting point or range of the total composition of the lipid phase does not exceed 25 °C, and more preferably is within the temperature ranges mentioned above.

[0131] In a particularly preferred embodiment, the products of this invention have a lipid phase containing:

- (a) at least 1 - 50 % (w/w), in particular at least 1 - 10 % of an oily or waxy component selected from C₁₄-C₃₀-dialkyl ethers, C₁₄-C₃₀-dialkyl carbonates, C₄-C₃₄-dicarbonic acids or C₁₂-C₃₀-hydroxyfatty alcohols or mixtures thereof
- (b) 0,1 - 5 % (w/w) of at least one active ingredient
- (c) 1 - 10 % (w/w) of at least one oil
- (d) 0.1 - 10 % (w/w) of at least one emulsifier
- (e) 5 - 90 % (w/w) of further waxy components
- (f) 0 - 5 % (w/w) water

Application of the lipid phase

[0132] The lipid phase may be applied to the sheet in various ways. Preferably the lipid phase is applied at the surface or at the surface portion of the sheet, on one or on both sides.

[0133] The lipid phase can be applied evenly or non-evenly to the sheet, non-evenly meaning that the distribution of the amount of the lipid phase varies over the area of the sheet, i.e. some areas of the sheet can have greater or lesser amounts of the lipid phase. Preferably the lipid phase is evenly applied to the area of the sheet.

[0134] The lipid phase can be applied discontinuously or continuously to one or both sides of the fabric, or it may even be applied as a complete covering of one or both surfaces of the fabric.

[0135] The lipid phase preferably is applied in a discontinuous pattern, to one or both sides of the sheet. To this purpose the lipid phase is applied in a predetermined, controlled manner to specific areas of the sheet. A discontinuous pattern is one in which the lipid phase has been applied to distinct regions separated by regions of the sheet which are free of the lipid phase. The lipid phase in that instance is applied to defined parts or regions of the sheet which may take a variety of forms. The lipid phase may in particular be applied as described above more generally for the application of both

phases. Particular for in which the lipid phase may be applied are, e.g. stripes, dots or spots, geometric configurations, either of regular or irregular shape, for example circles, ellipses, squares, rectangles and the like, logos, text, letters or any other noncontinuous pattern, including the patterns described hereinabove more generally for the application of the lipid and aqueous phase.

[0136] Discontinuous patterns also comprise essentially networks of larger patterns of the lipid phase. In a preferred embodiment, the lipid phase is present as discrete stripes which can be disposed discontinuously, i. e. interrupted, or preferably continuous over the whole surface of the wipe. The stripes may also form a pattern of discrete segments which collectively comprise a stripe or they may have a repetitive pattern such as a sinusoidal shape or wave-like and the like pattern. If waving stripes are selected, preferably the stripes are in phase, so that parallelism is maintained and each stripe remains equally spaced from the adjacent stripes.

[0137] The stripes are preferably oriented in the machine direction, for ease of manufacture.

[0138] The lipid phase may be colorless or colored, i. e. mono- or multi-colored. Multi-colored patterns are obtained by applying several lipid phases that have been dyed differently. If colored it will provide to alert the user of the fact that the sheet is covered by a special material that contains an active ingredient or it may also make the product aesthetically attractive.

[0139] In a particular embodiment, not more than half of the surface of the sheet, either on one side or, which is preferred, on both sides is carrying or covered by the lipid phase. In a preferred embodiment, the lipid phase is present at the surface on both sides, covering not more than 50 % of the sheet's surface, in particular covering not more than 35 % or not more than 25 % of the surface. In a particularly preferred embodiment, the lipid phase is present as stripes, in particular as parallel stripes running in parallel with the side of the sheet, covering not more than half or, more in particular 25 % of the surface. In another particularly preferred embodiment, the lipid phase is present as dots, equally spread over the entire surface of the sheet, covering not more than 50 % of the surface.

[0140] In case of stripes, the width of which preferably is between 1 to 10 mm, more preferably between 3 to 7 mm. In case of dots, round shapes are preferred, circles or ellipsoids, with an average diameter between 1 to 10 mm, more preferably between 3 to 7 mm.

[0141] The lipid phase is typically applied in an amount of from about 3 to 40 g/m², preferably from about 10 to about 20 g/m², either on one side or, preferably, on both sides of the sheet. Or, alternatively, the lipid phase is applied in an amount of about 0.06 g to 0.8 g per gram of substrate, preferably from about 0.20 g to 0.40 g per gram of dry substrate.

[0142] The lipid phase can be applied to the sheet by any method that can be used to contact or impregnate a liquid or molten lipid material to or in a sheet, in par-

particular any such method that allows coating of the lipid material onto the surface of the sheet. As used herein the term 'coating' refers to printing, covering, overlaying, finishing, spraying, extruding, laminating or any other method of applying the phase to the surface of the sheet. A particular coating technique is extrusion whereby the composition is forced through tubes in contact with the sheet whilst the sheet passes across the tube. A preferred technique comprises contacting the sheets with a heated head equipped with a slit blade, i.e. a blade having cut-out areas, wherefrom the lipid phase, in molten state, is extruded. Another preferred coating technique involves the so-called hot melt process which comprises spraying the liquefied lipid phase from a heated spraying head or nozzle. Another application technique involves spraying the composition on a rotating surface such as calender roll that then transfers the composition to the surface of the substrate. Still another technique is based on traditional printing technologies whereby a rotating surface is provided with elevations (by engraving, embossing or similar techniques) and the elevations are contacted with the liquefied lipid phase, e.g. by running it through a bath with liquefied phase one, and thus printed on the sheet. The lipid phase may also be applied by a combination of these application techniques.

[0143] The lipid phase preferably is applied in liquid form, e.g. in its molten form, or can also be applied as a powder.

[0144] The lipid phase can be applied either to one surface of the sheet or both surfaces, preferably to both surfaces.

[0145] The lipid phase preferably is applied in such manner that it will remain on the fabric surface during the manufacturing process and storage. This can best be accomplished by applying the lipid phase above its melting temperature, e.g. by spraying or coating it when molten to the surface of the sheet and subsequently allowing it to cool below its melting point so that the phase solidifies.

[0146] The lipid phase preferably is applied such that it is present at the surface of the sheet and because of its physical location in that instance, the lipid phase is readily available to be spread onto the skin during usage. As a result, the effectiveness with which the lipid phase is transferred to the skin during use, the availability and therefore the effectiveness of active ingredients is increased compared to products where the active agent is simply incorporated into a single continuously applied phase.

[0147] A melting point or range of above 25 °C, in particular within the ranges specified above, is preferred because it allows to apply the lipid phase in liquid (molten) state to the sheet, and subsequently, after it having been cooled, to be present in solid state on the sheet. Additionally this allows a more convenient and easy aftertreatment of the sheet to which the lipid phase has been applied in this manner, with the aqueous phase.

This allows the two phases to be applied in such manner that they do not mix or interact. The lipid phase forms a weak non-brittle film on the sheet. Sheets that have been treated this way are particularly stable, in particular during storing, essentially because mixing of the two phases is avoided in this way. Additionally such sheets will allow the lipid phase to melt upon contact with the skin, thus allowing a local mixing or emulsification of both phases.

The aqueous phase

[0148] The aqueous phase can be any of the art-known aqueous based formulations used to impregnate wipes. Beside water the aqueous phase may also contain further ingredients or additives such as surfactants, emulsifiers, consistency factors, conditioners, moisturizers, thickeners, preservatives, active ingredients, in particular dermatologically active ingredients, fragrances and the like. Active ingredients as mentioned herein comprise, for example, anti-inflammatories, antibacterias, anti-fungus and the like agents. Active ingredients suited for topical applications are particularly preferred.

[0149] The aqueous phase may further contain one or more **preservatives** such as, for example, phenox-yethanol, C₁₋₄ alkylparabens and their salts, in particular their alkali metal salts such as sodium salts (e.g. C₁₋₆ alkyl parabens such as methyl, ethyl, propyl, isopropyl, butyl paraben and the like parabens), chlorohexidine, formaldehyde or formaldehyde releaser, benzyl alcohol, chloroxylenol, phenoxyethanol, methylchloroisothiazolinone, methylisothiazolinone, sodium benzoate, chlorohexidine digluconate, methylidibromo glutaronitrile, sodium borate, 5-bromo-5-nitro-1,3-dioxane, alcohol, benzoic acid, dehydroacetic acid, diazolidinyl urea, dichlorobenzyl alcohol, glucose oxidase, hexamidine diisethionate, imidazolidinyl urea, iodopropynyl butylcarbamate, isobutylparaben, isopropylparaben, lactoperoxidase, magnesium nitrate, PEG-4 laurate, phenethyl alcohol, polyaminopropyl biguanide, potassium sorbate, propylene glycol, pyridoxine HCl, quaternium-15, sorbic acid, triclosan, tocopherol and the like.

[0150] Suitable **surfactants** for the aqueous phase comprise:

alkyl sulfates e.g. sodium lauryl sulfates, ammonium lauryl sulfates, sodium cetearyl sulfates, alkyl sulfoacetates e.g. sodium lauryl sulfoacetate, alkyl ether sulfates e.g. sodium laureth sulfates, sodium trideceth sulfates, sodium oleyl sulfates, ammonium laureth sulfates, alkyl ether sulfosuccinates e.g. disodium laureth sulfosuccinate, alkyl glycosides e.g. decyl glucoside, lauryl glucoside, alkyl isethionates, amphoterics e.g. cocamidopropyl betaine, sodium

cocoamphoacetate, sodium lauroamphoacetate, disodium lauroamphodiacetate, disodium cocoamphodiacetate, sodium lauroamphopropionate, disodium lauroamphodipropionate, potassium or ammonium salts of the aforementioned amphoterics, capryl/capramidopropyl betaine, andecylenamidopropyl betaine, lauramidopropyl betaine and fatty alcohol polyglycol ethers.

[0151] Suitable **consistency factors** are e.g. fatty alcohols or their mixtures with fatty acid esters, e.g. acetylated lanolin alcohol, aluminum stearates, carbomer, cetyl alcohol, glyceryl oleate, glyceryl stearate, glyceryl stearates (and) PEG 100 stearate, magnesium stearate, magnesium sulfates, oleic acid, stearic acid, stearyl alcohol, myristyl myristate, isopropyl palmitate, beeswax and synthetic equivalents thereof, carbomers, and the like, including mixtures thereof.

[0152] Suitable **conditioners** are e.g. alkylamido ammonium lactate, cetrimonium chloride and distearoylethyl hydroxyethylmonium methosulfates and cetearyl alcohol, cetyl dimethicone, cetyl ricinoleate, dimethicone, laureth-23, laureth-4, polydecene, and retinyl palmitate, agents selected from glyceryl monooleate and cocogluco-side including mixtures thereof (in particular the product 'Lamesoft ®' of Cognis which is a mixture of these two components), quaternised protein hydrolysates, quaternised cellulose and starch derivatives, quaternised copolymers of acrylic or methacrylic acid or salts, quaternised silicone derivatives, silicone oils, cyclomethicones, and the like agents, including mixtures thereof.

[0153] Suitable **thickeners** are e.g. acrylates/stearth-20 methacrylate copolymer, carbomer, carboxymethyl starch, cera alba, dimethicone/vinyl dimethicone crosspolymer, propylene glycol alginate, hydroxyethylcellulose, hydroxypropyl methylcellulose, silica, silica dimethyl silylate, xanthan gum, hydrogenated butylene/ethylene/styrene copolymer.

[0154] The aqueous phase may further comprise **film-forming substances** like chitosan and derivatives thereof, derivatives of poly acrylic acid, polyvinyl pyrrolidone and its derivatives, and the like.

Application of the aqueous phase

[0155] The aqueous phase may be applied to the sheet using methods generally known in the art for applying aqueous liquid lotions such as spraying, dripping, immersing and the like techniques. A preferred application method for the aqueous phase is by spraying with a suitable nozzle or by dripping, for example by using a perforated tube with holes or slits in it. The immersing technique can be done by running the sheets through a bath holding the lipid phase and subsequently controlling the amount of liquid that is absorbed by pressing.

[0156] The aqueous phase may be applied in various ways as described for the lipid phase, evenly or non-evenly, continuously or non continuously, at the surface

or surface portion or, preferably, throughout the whole of the sheet material. Optionally some parts of the sheet can be left dry, i.e. not having the lipid and the aqueous phase, or some parts can only have the lipid or the aqueous phase.

[0157] The aqueous composition is typically applied in an amount of from about 1.0 g to 10 g per gram of substrate, preferably from 2.0 g to 5 g per gram of substrate, most preferably from 2 g to 4.5 g per gram of dry substrate, most preferably about 3.7 to about 3.8 g per gram substrate. Or, the aqueous phase is applied in an amount of about 4 to about 8 g per wipe sized 17.2 x 21 cm, most preferably about 6 g per wipe.

[0158] The lotion comprising the aqueous phase may be applied on the entire area of the sheet, either throughout the whole of the sheet or only at its surface portions. In the latter instance the aqueous phase may be applied at both sides or only at one side of the sheet.

[0159] It may also be advantageous to only apply the lotion comprising the aqueous phase to only those areas (or that side) of the fabric which have (or has) not already been covered with the lipid phase. Furthermore it may also be advantageous to also leave certain areas of the sheet dry.

[0160] Since in many cases the product is used as a cleansing article it is useful to use the aqueous phase as cleanser. Soils that are most difficult to clean are either water insoluble and/or strongly adhere to the skin. Therefore the liquid used as the aqueous phase is formulated such that it is capable of taking up water-insoluble materials.

[0161] This can be achieved by two different approaches. Firstly the use of surface-active ingredients in an aqueous solution. Ingredients used in such formulations are polysorbate 20, cocamidopropyl betaine, disodium cocoamphodiacetate, sodium cocoamphoacetate, glyceryl stearate, cetareth-20, pareth-7 etc.. Secondly the use of an oil-in-water emulsion where the emulsification process of the soil is achieved via the uptake of thereof into the oil droplets. In both cases it is the effectiveness of the emulsification process, which governs the cleansing properties of the liquid used for impregnation.

Additional ingredients for either one or both phases

[0162] The lipid and/or the aqueous phase may contain further ingredients that may be present in one or in both phases.

Active ingredients

[0163] The lipid and/or the aqueous phase further may contain active ingredients for application to the skin. The lipid phase preferably contains oil-soluble or hydrophobic active agents, while the aqueous phase preferably contains water-soluble or hydrophilic active agents. However by using suitable emulsifiers oil-solu-

ble or lipophilic active ingredients can be incorporated into the aqueous phase and vice versa, water-soluble or hydrophilic agents can be incorporated in the lipid phase.

[0164] Products having a lipid and/or an aqueous phase that contains one or more active ingredients constitute particularly attractive embodiments of the present invention. Particularly preferred embodiments are those wherein the active ingredients are present in the lipid phase.

[0165] The active ingredients can be present in particular combinations.

[0166] Examples of active agents which may be hydrophobic or hydrophilic for use in the products of the invention comprise anti-microbials, e.g. anti-bacterials and antifungals, anti-inflammatory agents, anti-irritating compounds, anti-itching agents, moisturising agents, skin caring ingredients, plant extracts, vitamins, and the like. Examples of such ingredients comprise complexes of PVP and hydrogen peroxides, diclofenac, acetyl salicylic acid, salicylates, ibuprofen, bisabolol, mimosa extract (*Mimosa tenuiflora*), hyaluronic acid, propylene glycol, glycerines, chondroitin sulfates, plant extracts, bisabolol, panthenol, tocopherol, actives for anti-stinging, anti-irritants, anti-dandruffs, for anti-ageing e.g. retinol, melibiose etc. Other suitable actives are e.g. *medicago officinalis*, *actinidia chinensis*, allantoin, *Aloe barbadensis*, *anona cherimolia*, *anthemis nobilis*, *arachis hypogaea*, *arnica montana*, *avena sativa*, beta-carotene, bisabolol, *borago officinalis*, butylene glycol, *calendula officinalis*, *camellia sinensis*, camphor, *candida bombicola*, capryloyl glycine, *carica papaya*, *centaurea cyanus*, cetylpyridinium chloride, *Chamomilla recutita*, *chenopodium quinoa*, *chinchona succirubra*, *chondrus crispus*, *Citrus aurantium dulcis*, *Citrus grandis*, *Citrus limonum*, *Coco-nucifera*, *Coffea arabica*, *Crataegus monogina*, *Cucumis melo*, dichlorophenyl imidazolidioxolan, *enteromorpha compressa*, *equisetum arvense*, ethoxydiglycol, ethyl panthenol, farnesol, ferulic acid, *fragaria chiloensis*, *gentiana lutea*, *Ginkgo biloba*, glycerine, glyceryl laurate, *glycyrrhiza glabra*, *hamamelis virginiana*, heliotropine, hydrogenated palm glycerides, Citratess, hydrolyzed castor oil, hydrolyzed wheat protein, *hypericum perforatum*, *Iris florentina*, *juniperus communis*, lactic proteinum, lactose, *lawsonia inermis*, linalool, *linum usitatissimum*, lysine, magnesium aspartate, *magnifera indica*, *malva sylvestris*, mannitol, mel, *melaleuca alternifolia*, *mentha piperita*, menthol, menthyl lactate, *Mimosa tenuiflora*, *Nymphaea alba*, olafur, *oryza sativa*, panthenol, paraffinum liquidum, PEG-20M, PEG-26 Jojoba acid, PEG-26 Jojoba alcohol, PEG-35 castor oil, PEG-40 hydrogenated castor oil, PEG-60 hydrogenated castor oil, PEG-8 caprylic/capric acid, *persea gratissima*, petrolatum, potassium aspartate, potassium sorbate, propylene glycol, *prunus amygdalus dulcis*, *prunus armeniaca*, *prunus persica*, retinyl palmitate, *ricinus communis*, *rosa canina*, *rosmarinus officinalis*, *rubus idaeus*, salicylic acid, *sambucus nigra*,

sarcosine, *serenoa serrulata*, *simmondsia chinensis*, sodium carboxymethyl betaglucan, sodium cocoyl amino acids, sodium hyaluronate, sodium palmitoyl proline, stearoxytrimethylsilane, stearyl alcohol, sulfurized TEA-ricinoleate, talc, *Thymus vulgaris*, *tilia cordata*, tocopherol, tocopheryl acetate, trideceth-9, *triticum vulgare*, tyrosine, andecylenoyl glycine, urea, *vaccinium myrtillus*, valine, zinc oxides, zinc sulfates.

[0167] Of particular interest are active ingredients, that can be used for treating skin that shows inflammatory reactions, that is irritated, red or damaged. Examples of such agents are zinc compounds or sulphur.

[0168] The active ingredients can be present, depending on the nature of the ingredients and their application, in various concentrations, but usually are present in a quantity in the range of 0.01 - 10 % (w/w), preferably from 0.1 - 7 % (w/w) and more preferably 1 - 5 % (w/w).

Typical examples of **anti-microbial agents** are those active against gram-positive bacteria such as 2,4,4'-trichloro-2'-hydroxydiphenyl ether, chlorohexidine (1,6-di-(4-chlorophenyl-biguanido)hexan) or TCC (3,4,4'-trichlorocarbanilide). Furthermore many odorants and etheric oils have anti-microbial activity. Typical examples are the active ingredients eugenol, menthol and thymol in clove, mint and thyme oil. An interesting natural deodorizing agent is the terpene alcohol farnesol (3,7,11-trimethyl-2,6,10-dodecatrien-1-ol). Also glycerine monolaurate, glycerine stearate, glycerine oleate as well as glycerine dioleate have been found to possess anti-microbial activity and are particularly attractive for use in products that are applied on babies because of their mildness and harmlessness.

[0169] **Biogenic active ingredients** are for example tocopherol, tocopherolacetate, tocopherol palmitate, ascorbic acid, desoxyribonucleinic acid, retinol, bisabolol, allantoin, phytantriol, panthenol, α -hydroxycarbonic acids, amino acids, ceramides, pseudoceramides, essential oils, extracts and vitamin complexes. Preferred active ingredients are oil-soluble vitamins and vitamin predecessors. Particularly preferred are tocopherol (vitamin E) and tocopherol derivatives.

[0170] The quantity of anti-microbial agents can vary but usually is in the range of about 0.1 to 2 % (w/w) - relative to the total amount of the lipid and/or the aqueous phase. Glycerine esters can be used in larger quantities (see above).

Moisturizers

[0171] The lipid and/or aqueous phase can further contain one or more moisturizers. These are added to improve the sensoric properties as well as to regulate skin hydration. These agents additionally can improve the penetration of the composition in or into the sheet, in particular in or into the wipes.

[0172] Moisturizers typically are present in quantities of 1 - 20 % (w/w), preferably of 5 - 15 % (w/w), and more

preferably 5 -10 % (w/w).

[0173] Suitable moisturizers are a.o. amino acids, pyrrolidone carbonic acid, lactic acid and its salts, lactitol, urea and urea derivatives, ureic acid, glucosamine, creatinine, crack products of collagen, chitosan or chitosan salts/-derivatives, and in particular polyols and polyol derivatives (e.g. ethylene glycol, propylene glycol, butylene glycol, pentylene glycol, hexylene glycol, erythrite, 1,2,6-hexanetriol, polyethylene glycols such as PEG-4, PEG-6, PEG-7, PEG-8, PEG-9, PEG-10, PEG-12, PEG-14, PEG-16, PEG-18, PEG-20, PEG-135, PEG 150), sugar and sugar derivatives (a.o. fructose, glucose, maltose, maltitol, mannite, inositol, sorbitol, sorbitylsilylandiol, sucrose, trehalose, xylose, xylitol, glucuronic acid and its salts), ethoxylated sorbit (Sorbeth-6, Sorbeth-20, Sorbeth-30, Sorbeth-40), honey and hydrogenated honey, hydrogenated starch hydrolysates, as well as mixtures of hydrogenated wheat protein, hydrolyzed milk protein, lecithin, pythantriol, hyaluronic acid and salts thereof, and PEG-20-acetate copolymers. Particularly preferred moisturizers are glycerine, diglycerine and triglycerine.

[0174] The products according to this invention can be used as **anti-perspirants or deodorants**, in particular as wipes or tissues for use in these applications. In products for these applications either one or both phases contain actives that have deodorizing and /or anti-perspirant properties. Actives that can be used to this purpose are anti-perspirant agents such as, for example, aluminium chlorohydrates, aluminium-zirconium-chlorohydrate as well as zinc salts. Other such agents comprise aluminium hydroxylactates as well as acid aluminium/zirconium salts. A particularly suitable chlorohydrate is the compound of formula $[Al_2(OH)_5Cl] \cdot 2.5 H_2O$. Further such agents are aluminium-zirconium-tetrachlorohydroxy-glycine-complexes. Esterase inhibitors can be added as further deodorizing agents, i.e. agents such as trialkyl citrates such as trimethylcitrate, tripropyl citrate, triisopropyl citrate, tributyl citrate and in particular triethyl citrate. Further esterase inhibitors are sterol sulfates or -phosphates, such as, for example, lanosterine-, cholesterol-, campesterine-, stigmasterine- and sitosterinsulfate respectively-phosphate, dicarbonic acids and its esters, such as, for example, glutaric acid, glutaric acid monoethylester, glutaric acid diethylester, adipic acid, adipic acid monoethylester, adipic acid diethylester, malonic acid and malonic acid diethylester, hydroxycarbonic acids and their ester such as, for example, citric acid, malonic acid, tartaric acid or tartaric acid diethylester.

[0175] Antibacterial active ingredients, that influence the growing conditions and eradicate perspiration decomposing bacteria, or impede their growth, can also be present in the lipid and/or aqueous phase. Examples of such ingredients are chitosan, phenoxyethanol and chlorohexidine gluconate and in particular 5-chloro-2-(2,4-dichlorophenoxy)-phenol.

[0176] The products according to the invention can al-

so be used in sunscreen applications and in that instance take the form of sunscreen wipes. In these products the lipid and/or aqueous phase contains one or more **sunscreen filters** which are for example organic substances that are capable of absorbing ultraviolet radiation and to set free the absorbed energy as longer-wave radiation, e.g. as thermic energy. UVB-filters can be oil or water soluble. As oil-soluble substances there can be mentioned for example:

- 3-Benzylidene campher respectively 3-benzylidene norcampher and derivatives thereof, e.g. 3-(4-methylbenzylidene) campher;
- 4-Aminobenzoic acid derivatives, respectively 4-(dimethylamino)benzoic acid-2-ethylhexyl esters, 4-(dimethylamino)benzoic acid-2-octyl esters and 4-(dimethylamino)benzoic acid amylesters;
- Esters of cinnamonic acid, in particular 4-methoxycinnamonic acid-2-ethylhexylester, 4-methoxycinnamonic acid propylester, 4-methoxycinnamonic acid isoamyl ester, 2-cyano-3,3-phenylcinnamonic acid-2-ethylhexyl ester (octocrylene);
- Esters of salicylic acid, respectively salicylic acid-2-ethylhexylester, salicylic acid-4-isopropylbenzyl ester, salicylic acid homomenthyl ester;
- Derivatives of benzophenones, in particular 2-hydroxy-4-methoxybenzophenone, 2-hydroxy-4-methoxy-4'-methylbenzophenone, 2,2'-dihydroxy-4-methoxybenzophenone;
- Esters of benzalmalonic acid, in particular 4-methoxybenzmalonic acid di-2-ethylhexyl ester;
- Triazine derivatives, such as, for example, 2,4,6-trianilino-(p-carbo-2'-ethyl-1'-hexyloxy)-1,3,5-triazin and octyltriazone;
- Propane-1,3-diones, such as for example, 1-(4-tert-butylphenyl)-3-(4'-methoxyphenyl)propane-1,3-dione;
- Ketotricyclo(5.2.1.0)decane-derivatives.

Water-soluble UV-filter are for example:

[0177]

- 2-Phenylbenzimidazol-5-sulfonic acid and its alkali-, alkaline earth-, ammonium-, alkylammonium-, alkanolammonium- and glucammonium salts;
- Sulfonic acid derivatives of benzophenones, in particular 2-hydroxy-4-methoxybenzophenone-5-sulfonic acid and its salts;
- Sulfonic acid derivatives of 3-benzylidene campher, e.g. 4-(2-oxo-3-benzylidene methyl)benzol-sulfonic acid and 2-methyl-5-(2-oxo-3-benzylidene)sulfonic acid and its salts.

[0178] Typical UV-A-Filters that can be used are derivatives of benzoylmethane, such as, for example, 1-(4'-tert-butylphenyl)-3-(4'-methoxyphenyl)propane-1,3-dione, 4-tert-butyl-4'-methoxydibenzoylmethane

(Parsol 1789), or 1-phenyl-3-(4'-isopropylphenyl)-propane-1,3-dione. Mixtures of UV-A and UV-B-filters can be evidently used too.

[0179] Apart from the above mentioned soluble substances, there can also be used insoluble sunscreen pigments, namely finely dispersed metal oxides or metal salts. Examples of appropriate metal oxides are in particular zinc oxide and titanium dioxide as well as oxides of iron, zirconium, silicon, mangan, aluminium and cerium as well as mixtures thereof. Salts that can be used comprise silicates (talcum), barium sulfate or zinc stearate. The particle size of these pigments is sufficiently small, e.g. less than 100 nm, in particular between 5 and 50 nm and more in particular between 15 and 30 nm. The particles can be spherical but can have other shapes too such as ellipsoidal or similar shapes. The surface of the pigments may have been treated, e.g. hydrophilized or made hydrophobic. Typical examples are coated titanium dioxide, e.g. Titanium dioxide T 805 (Degussa) or Eusolex® T 2000 (Merck). Silicones can be used as hydrophobic coating agents, in particular trialkoxyoctyl silanes or simethicones. So-called micro- or nanopigments are particularly attractive for use in sunscreen products.

[0180] Apart from both groups of primary light protecting filters that are mentioned above there can also be used secondary light protecting factors. These pertain to the class of anti-oxidants and their activity is based on the interruption or decrease of the photochemical processes caused by solar radiation upon penetration in the skin.

[0181] Typical examples of secondary light protecting agents are amino acids such as for example glycine, histidine, tyrosine and tryptophane, including derivatives of amino acids; imidazoles (for example urocanic acid) and derivatives thereof; peptides such as D,L-carnosine, D-carnosine, L-carnosine and derivatives thereof (e.g. anserin). Further agents that can be used are carotinoides, carotenes (for example α -carotene, β -Carotene and lycopene) and derivatives thereof; chlorogenic acid and its derivatives; lipic acid and derivatives thereof (e.g. dihydrolipoic acid), aurothioglucose, propylthiouracil and other thioles (for example thioredoxine, glutathione, cysteine, cystamine and glycosyl-, N-acetyl-, methyl-, ethyl-, propyl-, amyl-, butyl- and lauryl-, palmitoyl-, oleyl-, γ -linoleyl-, cholesteryl- and glyceryl esters) and their salts. Further examples are dilauryl thiodipropionate, distearyl thiopropionate, thiodipropionic acids and derivatives thereof (for example esters, ethers, peptides, lipids, nucleotides, nucleosides and salts), sulfoximine compounds (for example buthionin sulfoximine, homocystein sulfoximine, butionine sulfone, penta-hexa-, and heptathione sulfoximine). These secondary agents usually are formulated into very low concentrations (e.g. pmol to μ mol/kg),

[0182] Other secondary agents (usually in small concentrations, as mentioned above) are chelating agents (for example α -hydroxy fatty acids, palmeate acid,

phytic acid, lactoferrin), α -hydroxy acids (for example citric acid, lactic acid, malic acid), humic acid, bile acid, bile extracts, bilirubin, biliverdin, EDTA, EGTA and derivatives thereof; unsaturated fatty acids and derivatives thereof such as for example γ -linolenic acid, linoleic acid, oleic acid, folic acid and derivatives thereof, ubiquinones and ubiquinol and derivatives thereof, vitamin C and derivatives thereof (e.g. ascorbyl palmitate, Mg ascorbylphosphate, ascorbyl acetate), tocopherol and derivatives thereof (for example vitamin E acetate), vitamin A and derivatives thereof (e.g. vitamin A palmitate), coniferyl benzoates of benzoic acid, rutinic acid and derivatives thereof, α -glycosyl rutin, ferula acid, furfurylidene glucitol, carnosine, butyl hydroxytoluene, butyl hydroxyanisol, nordihydroguaiac resin acid, nordihydroguaiaretic acid, trihydroxybutyrophenone, ureic acid and derivatives thereof, mannose and derivatives thereof, superoxide-dismutase, zinc and derivatives thereof such as zinc oxide, zinc sulphate, selenium and derivatives thereof (e.g. seleno-methionine); stilbene and derivatives thereof (e.g. stilbene oxide, trans stilbene oxide); and any appropriate derivatives of these UV filters.

[0183] To improve the rheological behavior there can be added **hydrotropes**, such as, for example, ethanol, isopropyl alcohol, or polyols. Polyols, that can be used in particular have 2 to 15 carbon atoms and at least 2 hydroxyl groups, and optionally have further substituents such as amino or other nitrogen-based substituents. A number of these compounds have been mentioned amongst the moisturizing agents. Typical examples are:

- glycerine;
- alkylene glycoles, such as, for example, ethylene glycol, diethylene glycol, propylene glycol, butylene glycol, hexylene glycol as well as polyethylene glycol with a mean molecular weight from 100 to 1.000 Daltons;
- technical oligoglycerine mixtures with a condensation level of 1,5 to 10 such as technical diglycerine mixtures with a diglycerine content of 40 to 50 % (w/w);
- methylol compounds such as in particular trimethylol ethane, trimethylol propane, trimethylol butane, pentaerythrite and dipentaerythrite;
- lower alkyl glucosides, in particular those with 1 to 8 carbon atoms in the alkyl rest, for example methyl- and butyl glucoside;
- sugar alcohols with 5 to 12 carbon atoms, such as, for example, sorbit or mannit,
- sugars with 5 to 12 carbon atoms, such as, for example, glucose or saccharose;
- amino sugars, such as, for example, glucamine;
- dialcohol amines, such as diethanolamine or 2-amino-1,3-propane diol.

[0184] As **self-tanning agents** there can be added

dihydroxy acetone.

[0185] As **perfume oils** there can be mentioned mixtures from synthetic or natural odorous substances. Natural odorous substances are extracts from blossoms (lily, lavender, rose, jasmine, neroli, ylang-ylang), from stems and leaves (geranium, patchouli, petitgrain), from fruits (anis, coriander, caraway, juniper), from cortex (bergamot, lemon, orange), from roots (macis, angelic, celery, cardamon, costus, iris, calmus), from wood (pine, sandelwood, guajak, cedar, rose), from herbs and grass (tarragon, lemongrass, sage, thyme), from needles and branches (spruce, fir, pine, mountain pine), from resins and balms (galbanum, elemi, benzoin, myrrh, olibanum, opoponax). Furthermore animal odorous substances can be used, such as, for example, civet and castoreum. Typical synthetic odorous substances are products of the classes of esters, ethers, aldehydes, ketones, alcohols and hydrocarbons.

Etheric oils of lower volatility, which are mostly used as aromatic components, can be added to perform as perfume oils, e.g. sage, camomile, clove, balm, mint, cinnamon leaf, lime blossom, juniper, vetiver, olibanum, galbanum, labolanum and lavandin. Particular oils are bergamot, dihydro-myrcenol, lillial, lyral, citronellol, phenylethyl alcohol, α -hexylcinnamonic aldehyde, geraniol, benzyl acetone, cyclamene aldehyde, linalool, boisambrene forte, ambroxan, indole, hedione, sandelice, lemon, mandarin, orange, allyl amylglycolate, cyclovertal, lavandin, muskateller sage, β -damascone, geranium bourbon, cyclohexyl salicylate, vertofix coeur, iso-E-super, fixolide NP, evernyl, iraldein gamma, phenyl acetic acid, geranyl acetate, benzyl acetate, rose oxide, romillat, irotyl and floramate; or mixtures thereof.

[0186] The lipid and/or aqueous phase may contain cosmetically acceptable **dyes** which can be present in quantities in the range of 0,001 to 0,1 % (w/w), relative to the total quantity of the lipid and/or aqueous phase. Oil soluble dyes preferably are used in the lipid phase, water-soluble dyes in the aqueous phase. Preferably, the lipid phase contains one or more dyes, the aqueous phase not. Dyes that can be used in the lipid phase are, for example the C.I. series of oil-soluble dyestuffs, e.g. C.I. 47000, C.I. 67565, C.I. 26100, C.I. 60725, C.I. 12150, C.I. 75810, C.I. 75300.

[0187] The addition of a dye has the advantage that it provides of a visible indication for the user, sending the message of particular (active) ingredients having been incorporated in the lipid phase. It allows furthermore to visualize the stability of the phase, in particular of the lipid phase, that has been applied on the sheet can be easily visualized. This allows, for example, to monitor whether the oily and aqueous phases have become mixed upon the storage.

Emulsifiers

[0188] The lipid and/or aqueous phase in the products of the invention may further contain one or more emul-

sifiers which can be of the W/O (for use in the lipid phase) or the O/W (for use in the aqueous phase) type. The addition of an emulsifier allows the incorporation of hydrophilic components or agents into the lipid phase and vice versa of lipophilic components or agents into the aqueous phase.

Preferred are non-ionic emulsifiers which typically have good skin compatibility. Improved sensoric properties are obtained when combining non-ionic W/O and O/W emulsifiers. The lipid and/or aqueous phase may contain the emulsifier(s) in an amount of 0 to 20 % (w/w), respectively 0.1 to 15 % (w/w) and in particular 0.1 to 10 % (w/w) relative to the total quantity of the lipid and/or aqueous phase.

Non-ionic emulsifiers

[0189] Particular non-ionic emulsifiers comprise:

- (1) Addition products of 2 to 50 moles of ethylene oxide and/or 0 to 20 moles propylene oxide to linear fatty alcohols having 8 to 40 C-atoms, to fatty acids with 12 to 40 C-atoms and to alkylphenols with 8 to 15 C-atoms in the alkyl rest.
- (2) C_{12/18}-fatty acid mono- and -diesters of addition products of 1 to 50 moles of ethylene oxide and glycerine.
- (3) Glycerine mono- and -diesters and sorbitan mono- and -diesters of saturated and unsaturated fatty acids with 6 to 22 C-atoms and their ethylene oxide addition products.
- (4) Alkyl mono- and -oligoglycosides with 8 to 22 C-atoms in the alkyl rest and their ethoxylated analogs.
- (5) Addition products of 7 to 60 moles of ethylene oxide to castor oil and/or hardened castor oil.
- (6) Polyol- and in particular polyglycerine esters, such as e.g. polyol poly-12-hydroxystearate, polyglycerine polyricinoleate, polyglycerine diisostearate or polyglycerine dimerate. Also applicable are mixtures of compounds of several of these substance classes.
- (7) Addition products of 2 to 15 moles of ethylene oxide to castor oil and/or hardened castor oil.
- (8) Partial esters derived from linear, branch chained, unsaturated or saturated C₆-C₂₂-fatty acids, ricinoleic acid as well as 12-hydroxystearic acid and glycerine, polyglycerine, pentaerythrit, dipentaerythrit, sugar alcohols (e.g. sorbitol), alkylglucosides (e.g. methylglucoside, butylglucoside, laurylglucoside) as well as polyglucosides (e.g. cellulose), or mixed esters such as e.g. glyceryl stearate/citrate and glyceryl stearate/lactate.
- (9) Wool wax alcohols.
- (10) Polysiloxane-polyalkyl-polyether-copolymers and derivatives thereof.
- (11) Mixed esters from pentaerythrit, fatty acids, citric acid and fatty alcohols and/or mixed esters of

fatty acids with 6 to 22 C-atoms with methylglucose and polyols, respectively glycerine or polyglycerine.

(12) Polyalkylene glycols.

[0190] The addition products of ethylene oxide and/or of propylene oxide and fatty alcohols, fatty acids, alkylphenols, glycerine mono- and -diesters as well as sorbitan mono- and -diesters of fatty acids or of castor oil are known and commercially available products. Usually these are mixtures of homologues of which the average degree of alkoxylation corresponds to the ratio of starting quantities of ethylene oxide and/or propylene oxide and substrate, with which the addition reaction is conducted. Depending upon the degree of alkoxylation these products are either W/O- or O/W-emulsifiers. C_{12/18}-fatty acid mono- and -diesters of addition products of ethylene oxide to glycerine are known as re-fattening agents in cosmetic applications.

[0191] Particular useful and mild emulsifiers are polyolpoly-12-hydroxystearates and mixtures thereof with other components, that are available under the trade-name "Dehymuls® PGPH" (W/O-emulsifier) or "Eumulgin® VL 75" (1:1 w/w mixture with coco-glucosides, O/W-emulsifier) or Dehymuls® SBL (W/O-Emulsifier) from Cognis Deutschland GmbH. The polyol components of these emulsifiers can be derived from materials that have at least two and in particular 3 to 12 and more in particular 3 to 8 hydroxyl groups, and 2 to 12 carbon atoms.

[0192] In case it is desirable to incorporate water-soluble active ingredients and/or small amounts of water into the lipid phase it can be advantageous to add an emulsifier selected from the group of non-ionic O/W-emulsifiers (HLB-value: 8 - 18) and/or solubilizers. These can for example be the already mentioned ethylene oxide-adducts with a corresponding high degree of ethoxylation e.g. 10 - 20 ethylene oxide units in the case of O/W-emulsifiers and 20 - 40 ethylene oxide units for so-called solubilizers. Particularly attractive as O/W emulsifiers are Cetareth-12 und PEG-20 stearate. Particularly attractive solubilizers are Eumulgin® HRE 40 (INCI: PEG-40 Hydrogenated Castor Oil), Eumulgin® HRE 60 (INCI: PEG-60 Hydrogenated Castor Oil), Eumulgin® L (INCI: PPG-1-PEG-9 Laurylglycoether) and Eumulgin® SML 20 (INCI: Polysorbate-20).

[0193] Non-ionic emulsifiers of the group of alkyl oligoglycoside are particularly skin-compatible and therefore preferred as O/W-Emulsifiers. C₈-C₂₂-alkyl mono- and - oligoglycosides, their preparation and used have been described in the prior art. Oligoglycosides are meant to comprise oligomeric glycosides with a degree of oligomerisation of up to about 8. The degree of oligomerisation can also be a statistical average used for those products comprised of a specific range of oligoglycosides. An example is the product sold under the trade-name Plantacare® which has a C₈-C₁₆-alkyl group glycosidically bound to an oligoglucoside rest, having an

average degree of oligomerisation between 1 and 2.

Other non-ionic emulsifiers are the acyl glucamides. Preferred is the product sold under the tradename Emulgate® PL 68/50 (Cognis Deutschland GmbH) which is a 1:1 - mixture of alkyl polyglucosides and fatty alcohols, and a mixture of lauryl glucoside, polyglyceryl-2-dipolyhydroxystearate, glycerine and water, sold under the trade name Eumulgin® VL 75.

[0194] Lipophilic W/O-emulsifiers in principle are emulsifiers with a HLB-value in the range of 1 to 8, that are described, for example, in Kirk-Othmer, "Encyclopedia of Chemical Technology", 3rd Ed., 1979, Vol. 8, p. 913. The HLB-value of ethoxylated products is calculated by the formula: $HLB = (100 - L) : 5$, wherein L is the percentage (in weight %) of lipophilic groups, i.e. of fatty alkyl- or fatty acyl groups in the ethylene oxide adducts.

[0195] Particularly attractive W/O-emulsifiers are the partial esters of polyols, in particular of mono-, di- or tri-, sesqui esters of fatty acids of polyols, more in particular of C₃-C₆-polyols, such as, for example, glyceryl monoesters, partial esters of pentaerythritol or carbohydrate esters, e.g. saccharose distearate, or sorbitane mono-, di-, tri- or sesqui fatty esters in particular stearates, oleates, erucates, ricinoleates, hydroxystearates, isostearates (but also: tartrates, citrates, maleates) and the like. Also attractive are addition products of 1 to 30, respectively 5 to 10 moles ethylene oxide to these sorbitane esters.

Further Surfactants/Emulsifiers for both phases

[0196] Depending upon the use of the products of the present invention, the lipid and/or aqueous phase may further contain zwitterionic, amphoteric, cationic and or anionic surfactants.

[0197] Zwitterionic surfactants are those tensioactive compounds, that contain at least a quaternary ammonium group and at least a -COO⁻ or -SO₃⁻ group. Particularly useful zwitterionic surfactants are the so-called betaines such as N-alkyl-N,N-dimethyl ammonium glycinate, for example coco-alkyl dimethylammonium glycinate, N-acylaminoethyl-N,N-dimethylammonium glycinate, for example coco-acyl aminoethyl dimethylammonium glycinate, and 2-alkyl-3-carboxymethyl-3-hydroxyethylimidazoline, each having 8 to 18 C-Atoms in the alkyl- or acyl group as well as coco-acyl amino ethyl hydroxyethyl carboxymethyl glycinate. A preferred zwitterionic surfactant is the fatty acid amide-derivative known by its INCI-name cocamidopropyl betaine.

[0198] Ampholytic surfactants can further be added, in particular as co-surfactants. Ampholytic surfactants are understood to comprise those tensioactive compounds, that beside a C₈-C₁₈-alkyl- or acyl group at least contain a free amino group and at least a -COOH- or -SO₃H- group and are able to form internal salts. Examples of appropriate ampholytic surfactants are N-alkyl glycines, N-alkyl propionic acids, N-alkyl amino bu-

teric acids, N-alkyl iminodipropionic acids, N-hydroxyethyl-N-alkyl amidopropyl glycines, N-alkyl taurine, N-alkyl sarcosine, 2-alkylaminopropionic acids and alkylamino acetic acids with in each alkyl group about 8 to 18 C-atoms.

[0199] Most preferred ampholytic surfactants N-coco-alkyl aminopropionate coco-acyl amino ethylamino propionate and C₁₂₋₁₈-acylsarcosine.

[0200] Anionic surfactants are characterized by a water solubilizing anionic group such as a carboxylate-, sulfate-, sulfonate- or phosphate- group and a lipophilic rest. Particular anionic surfactants are the alkali-, ammonium- or alkanol ammonium salts of alkyl sulfates, alkyl ethersulfates, alkyl ethercarboxylates, acyl isethionates, acyl sarkosinates, acyl taurines with linear alkyl- or acyl groups having 12 to 18 C-atoms as well as alkali- or ammonium salts of sulfosuccinates and acyl glutamates.

[0201] Quaternary ammonium derivatives can in particular be used as cationic surfactants. Preferred are ammonium halogenides, in particular chlorides and bromides, e.g. alkyl trimethylammonium chloride, dialkyl dimethylammonium chloride and trialkyl methylammonium chloride, z. B. cetyl trimethylammonium chloride, stearyl trimethylammonium chloride, distearyl dimethylammonium chloride, lauryl dimethylammonium chloride, lauryl dimethylbenzylammonium chloride and tri-cetyl methylammonium chloride. Additional cationic surfactants are the quaternary esters with good biological degradability, such as, for example, dialkylammonium methosulfates and methylhydroxyalkyl dialkoyloxy alkylammonium methosulfates (sold under the tradename Stepantex® and the products of the Dehyquart® -series). The term "Esterquats" is meant to comprise quaternized fatty acid triethanolamine ester salts which have a beneficial impact on the softness of the phases, in particular of the lipid phase. Further cationic surfactants are the quaternized protein hydrolysates.

Application and properties

[0202] The products according to the present invention advantageously result in an optimal release of the active ingredient(s), in particular when incorporated in the lipid phase, onto the skin during use.

[0203] Optimal release of active ingredients can be achieved by using a lipid phase which is a solid lipid having a melting point or melting range which is equal to or slightly exceeds body temperature. Without being bound to theory, it is believed that this results in a quicker melting of the lipid phase causing a faster and more efficient transfer and release to the skin of the active materials.

[0204] Optimal release of active ingredients can also be achieved by using a suitable emulsifier in one or both of the phases to cause a local emulsification process on the skin during use of the wipes. Preferably the emulsifier is present in the aqueous phase. This local emulsi-

fication may be the result of body temperature causing the lipid phase to melt or it may be the result of pressure exerted during usage of the wipe, or it may be the result of both, the latter being usually the case. In the instance of local emulsification by the effect of pressure, the emulsification process is driven by the (limited) pressure exerted by the user when applying the wipe, e.g. by rubbing it across the skin, dabbing it and the like. This causes the two phases to contact and form an emulsion locally.

[0205] In this local emulsification process, a limited amount of the phase without emulsifier is incorporated into the phase having the emulsifier. In a preferred embodiment, the aqueous phase contains a small amount of emulsifier, for example the emulsifier may be present in an amount from about 0.5 to about 5%, more in particular from about 1 to about 3%. In that instance some of the lipid phase is locally emulsified into the aqueous phase.

[0206] Although in preferred executions the lipid phase is not present on the whole surface of the wipe, good release of the lipid phase and of the components contained therein is attained, in particular when the local emulsification process comes into play.

[0207] Optimal release of active ingredients can also be achieved by making use of both above possibilities.

Manufacture.

[0208] This invention further concerns a process for preparing a product as defined herein, said process comprising contacting a porous or absorbent sheet with a lipid phase composition and an aqueous phase composition as described herein. The porous or absorbent sheet in particular is made of a non-woven material. The process comprises contacting the sheet simultaneously or subsequently with the lipid phase and the aqueous phase.

[0209] Contacting the sheet with the aqueous phase comprises impregnating it with the aqueous phase by procedures such as, for example, running through a bath, immersing, spraying, dripping and the like techniques.

[0210] Contacting the sheet with the lipid phase is as described above in the section 'lipid phase', preferably by spraying, printing or by a direct contact procedure in which there is a direct contact between the sheet and an application head having slit nozzles.

[0211] In a particular execution, the process comprises contacting the sheet with a lipid phase and subsequently with an aqueous phase.

[0212] The lipid and aqueous phases can be applied to the sheet at any time during the manufacturing process of the sheet, for example either one or both of the phases may be applied during the manufacturing process of the sheet material. Preferably the lipid and/or aqueous phase can be applied to the sheet after finishing the manufacturing process of the sheet, more pref-

erably after the sheet has been dried.

[0213] In a particular execution, the carrier material is cut into strips, the transversal size of which being similar to the size of the tissue, wipe or towelette. Subsequently, the lipid and aqueous phases are applied to these strips, preferably first the lipid phase and then the aqueous phase. Thereafter the strips are folded according to methods generally known and applied in the art. In an alternative execution, the lipid phase is applied to these strips, which are subsequently folded and the thus folded strips are moistened with the aqueous phase as, said moistening preferably comprising spraying or dripping, or by immersing in or running the strip through a bath containing the aqueous phase. The latter can also be sprayed or printed onto the strips.

[0214] In a further step, the strips are cut so that the desired size of the sheets, in particular of the wipes, is obtained. The thus obtained sheets (or wipes) can be packed individually or can be stacked in a determined number, e.g. a number between 10 and 30, preferably between 15 and 25, most preferably about 20, or a number between 50 and 100, preferably between 60 and 80, most preferably about 72, and the stack then packed in a suitable package, for example a plastic wrap, box and the like.

[0215] The products according to the invention can take the form of baby or adult wipes and can be used in a wide range of applications as personal care products, comprising, for example, baby cleansing wipes, face or body cleansing wipes, wipes for skin treatment or skin conditioning such as for example skin moisturization and against skin aging, insect repellent wipes, powder wipes, toilet wipes, anti-perspirant wipes, peeling wipes, after-sun treatment wipes, sunscreen wipes, wipes for feminine hygiene, and the like.

[0216] The products of the invention may find use as cleansing tools, however their use is not limited to this application only. They have been found to be more effective cleansers compared to products that have only an aqueous phase. This is due, i.a., by the fact that they can remove both aqueous and lipid soils and components.

[0217] The products described herein find use as applicators of active substances, in particular of the active substances mentioned herein, or they find use as both cleanser and applicator of active substances in one product.

[0218] Traditionally, wipes have been used primarily as a cleansing tool. Applications as a vehicle for active substances, i.e. so-called leave-on products, have been limited because of the poor transfer rate of the active ingredients from the wipe to the skin. The products of this invention provide a solution to this problem in that they result in an excellent transfer of active ingredients to the skin thus widening the applications of wipe products as a vehicle for a number of actives, in particular more expensive actives that so far could not be applied through wipes because of poor transfer rate. The prod-

ucts of this invention not only provide a more efficient transfer of active ingredients to the skin, but moreover provide other consumer benefits such as a more even distribution of the actives on the skin, better skin penetration.

[0219] The products of this invention show the additional advantage that they may combine in one and the same product both cleansing capability and the transfer of active ingredients to the skin, i.e. the application of leave-on products.

[0220] Either of both aspects may be present in a larger extend, i.e. the product may be primarily for cleansing purposes but also having the capability of transferring certain beneficial components or active substances to the skin, or vice versa, the products may be designed for applications in instances where the primary benefit is not cleansing but a better and more convenient form of application of leave-on products. Hence the products of this invention allow the possibility to be optimized against two key consumer needs - cleansing and caring.

[0221] The products of the invention therefore show improved performance in terms of cleansing and skin benefits since both benefits can be formulated in different phases independently.

[0222] Another benefit of the products of this invention is that they offer a softer feel of the fabric due to the modification of the fabric surface caused by the presence of the lipid phase. The products moreover offer gentler cleansing because of less friction of the wipe on the skin (softer skinfeel).

[0223] As used herein, softness refers to the tactile sensation perceived when the consumer holds the product, rubs it across the skin, or crumples it with the hand.

[0224] The products of this invention additionally offer the possibility to incorporate into or apply to one product two or more incompatible ingredients, thus allowing the user to apply incompatible agents with one and the same product. In particular it is possible to have a product that has as well water soluble as lipid soluble ingredients, for example a wipe that has both active ingredients that are water soluble and oil soluble.

[0225] A still further advantage lies in the fact that the instant products allow an improved transfer of actives onto the skin since the active ingredients are concentrated on the surface of the sheet / fabric material and not included in the inner phase of a typical o/w-emulsion.

[0226] The products according to the invention, and in particular the particular compositions of the lipid and aqueous phase described in this specification, possess the additional advantage that they are almost odorless (unless fragrances are added), environmentally friendly and biologically decomposable.

[0227] The products of this invention are particularly attractive because they allow convenient and quick application (easy to carry), and an easier and more evenly distribution of the product. They moreover are easy to apply on babies and children. The products are more efficient in that they allow faster cleansing.

[0228] In view of these beneficial properties, the products of this invention can be used in a wide variety of cosmetic and personal care applications, but also in other cleaning or cleansing applications such as cleaning of hard surfaces.

Examples

[0229] The following examples are given with the nomenclature of INCI. As used in the following examples, C.I. refers to dyes.

Example 1: lipid phases	
Phase 1-A	
Cocoglycerides	64.99 %
Cetyl Alcohol	33.00 %
Di-Stearyl Ether	1.00 %
Tocopherol	1.00 %
C.I. 61565	0.01 %
Phase 1-B	
Cocoglycerides	54.99 %
Cetyl Alcohol	33.00 %
Ceteareth-12	3.00 %
Glyceryl Stearate	4.00 %
Di-Stearyl Carbonate	2.00 %
Tocopherol	1.00 %
C.I. 61565	0.01 %
Aqua	2.00 %
Phase 1-C	
Cocoglycerides	49.99 %
Cetearyl Alcohol	20.00 %
Cegesoft® HF 52	5.00 %
Cegesoft® PS 6	3.00 %
Ceteareth-12	2.00 %
Glyceryl Stearate	2.00 %
PEG-20 Stearate	10.00 %
Di-Stearyl Ether	2.00 %
Tocopherol	1.00 %
C.I. 61565	0.01 %
Aqua	5.00 %
Phase 1-D	
Cocoglycerides	58.99 %
Glyceryl Stearate	25.00 %
Glyceryl Laurate	14.00 %
Di-Stearyl Carbonate	1.00 %
Tocopherol	1.00 %
C.I. 75300	0.01 %

(continued)

Example 1: lipid phases	
Phase 1-D	
Phase 1-E	
Cocoglycerides	30.00 %
Cetearyl Alcohol	1.00 %
Cegesoft® HF 52	20.00 %
Cegesoft® GPO	5.00 %
Ceteareth-12	15.00 %
Glyceryl Stearate	20.00 %
Di-Stearyl Ether	5.00 %
Tocopherol	1.00 %
Panthenol	1.00 %
Aqua	2.00 %
Phase 1-F	
Cocoglycerides	19.99 %
Cetearyl Alcohol	30.00 %
Cegesoft® PS 6	10.00 %
Eumulgin® VL 75	10.00 %
Ceteareth-12	5.00 %
Glyceryl Stearate	10.00 %
Di-Stearyl Carbonate	5.00 %
Tospearl® 145 A	5.00 %
Zinc Stearate	2.00 %
C.I. 61565	0.01 %
Aqua	3.00 %
Phase 1-G	
Myristyl Alcohol	19.99 %
Cocoglycerides	10.00 %
Cegesoft® HF 52	20.00 %
Eumulgin® VL 75	10.00 %
Glyceryl Stearate	20.00 %
PEG-20 Stearate	5.00 %
Di-Stearyl Carbonate	2.00 %
Panthenol	3.00 %
C.I. 61565	0.01 %
Aqua	10.00 %
Phase 1-H	
Myristyl Alcohol	47.99 %
Stearyl Alcohol	25.00 %
Eumulgin® VL 75	2.00 %
PEG-20 Stearate	14.00 %
1,2-Hexadecanediol	5.00 %
Bisabolol	1.00 %

(continued)

Example 1: lipid phases	
Phase 1-H	
C.I. 47000	0.01 %
Aqua	5.00 %
Phase 1-I	
Cocoglycerides	47.99 %
Stearyl Alcohol	20.00 %
Eumulgin® VL 75	2.00 %
PEG-20 Stearate	12.00 %
Di-Stearyl Carbonate	5.00 %
Cyclomethicone	3.00 %
Tospearl® 145 A	5.00 %
C.I. 75300	0.01 %
Aqua	5.00 %
Phase 1-J	
Cocoglycerides	55.99 %
Glyceryl Stearate	20.00 %
Glyceryl Laurate	15.00 %
Di-Stearyl Carbonate	5.00 %
Talc	2.00 %
Aluminum Starch Octenylsuccinate	2.00 %
C.I. 60725	0.01 %
Phase 1-K	
Cocoglycerides	50.99 %
Glyceryl Stearate	25.00 %
Glyceryl Laurate	15.00 %
Di-Stearyl Ether	5.00 %
Talc	2.00 %
Timiron® Splendid Gold	2.00 %
C.I. 21230	0.01 %
Phase 1-L	
Myristyl Alcohol	58.99 %
Stearyl Alcohol	23.00 %
PEG-20 Stearate	15.00 %
Di-Stearyl Carbonate	2.00 %
Panthenol	1.00 %
C.I. 61525	0.01 %
Phase 1-M	
Myristyl Alcohol	47.99 %
Stearyl Alcohol	25.00 %
Eumulgin® VL 75	2.00 %
PEG-20 Stearate	10.00 %

(continued)

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Example 1: lipid phases	
Phase 1-M	
Di-Stearyl Ether	7.00 %
Panthenol	2.00 %
C.I. 61525	0.01 %
Aqua	6.00 %
Phase 1-N	
Myristyl Alcohol	50.00 %
Stearyl Alcohol	25.00 %
Eumulgin® VL 75	2.00 %
PEG-20 Stearate	10.00 %
Di-Stearyl Ether	7.00 %
Ethyl Butylacetylaminopropionate	5.00 %
Panthenol	1.00 %
Phase 1-O	
Cocoglycerides	54.99 %
Cetyl Alcohol	33.00 %
Ceteareth-12	3.00 %
Glyceryl Stearate	4.00 %
Di-Stearyl Carbonate	2.00 %
Octyl Methoxycinnamate	6.00 %
C.I. 61565	0.01 %
Phase 1-P	
Cocoglycerides	56.99 %
Glyceryl Stearate	25.00 %
Glyceryl Laurate	14.00 %
Di-Stearyl Carbonate	1.00 %
Polyethylene	3.00 %
C.I. 75300	0.01 %
Phase 1-Q	
Cocoglycerides	58.93 %
Glyceryl Stearate	25.00 %
Glyceryl Laurate	15.00 %
Di-Stearyl Ether	1.00 %
Aqua	0.06 %
C.I. 61565	0.01 %
Phase 1-R	
Cocoglycerides	43.93 %
Stearyl Alcohol	15.00 %
Glyceryl Stearate	25.00 %
Glyceryl Laurate	15.00 %
Di-Stearyl Ether	1.00 %

(continued)

Example 1: lipid phases	
Phase 1-R	
Aqua	0.06 %
C.I. 61565	0.01 %
Phase 1-S	
Cocoglycerides	44.93 %
Glyceryl Stearate	25.00 %
Glyceryl Laurate	15.00 %
Di-Stearyl Ether	15.00 %
Aqua	0.06 %
C.I. 61565	0.01 %

Example 2: aqueous phases	
Phase 2-A	
Aqua	96.336 %
Polysorbate 20	0.600 %
PEG-75 Lanolin	0.100 %
Parfum	0.150 %
PEG-40 Hydrogenated Castor Oil	0.400 %
Propylene Glycol	1.120 %
Phenoxyethanol	0.800 %
Tetrasodium EDTA	0.078 %
Chamomilla Recutita	0.070 %
Ethoxydiglycol	0.171 %
Butylene Glycol	0.035 %
Glucose	0.016 %
Iodopropynyl Butylcarbamate	0.010 %
PEG- 4 Laurate	0.090 %
Citric Acid	0.020 %
Phase 2-B	
Aqua	98.252 %
Phenoxyethanol	0.800 %
Iodopropynyl Butylcarbamate	0.010 %
PEG- 4 Laurate	0.090 %
Parfum	0.150 %
Tetrasodium EDTA	0.078 %
Citric Acid	0.020 %
Polysorbate 20	0.600 %
Phase 2-C	
Aqua	97.250 %
Glycerines	1.000 %
Phenoxyethanol	0.800 %
Iodopropynyl Butylcarbamate	0.010 %

(continued)

Example 2: aqueous phases	
Phase 2-C	
PEG- 4 Laurate	0.090 %
Parfum	0.150 %
Tetrasodium EDTA	0.078 %
Citric Acid	0.020 %
Polysorbate 20	0.600 %
Phase 2-D	
Aqua	96.332 %
Glycerines	1.000 %
Phenoxyethanol	0.800 %
Polysorbate 20	0.600 %
PPG-15 Stearyl Ether	0.400 %
PEG-7 Glyceryl Cocoate	0.100 %
Propylene Glycol	0.350 %
Iodopropynyl Butylcarbamate	0.010 %
PEG- 4 Laurate	0.090 %
Chamomilla Recutita	0.070 %
Parfum	0.150 %
Tetrasodium EDTA	0.078 %
Citric Acid	0.020 %
Phase 2-E	
Aqua	97.33 %
Phenoxyethanol	0.800 %
Polysorbate 20	0.600 %
Sorbeth-30	0.400 %
Propylene Glycol	0.350 %
Dimethicone Copolyol	0.100 %
Iodopropynyl Butylcarbamate	0.010 %
PEG- 4 Laurate	0.090 %
Chamomilla Recutita	0.070 %
Parfum	0.150 %
Tetrasodium EDTA	0.078 %
Citric Acid	0.020 %
Phase 2-F	
Aqua	97.332 %
Phenoxyethanol	0.800 %
PEG-80 Sorbitan Laurate	0.600 %
Propylene Glycol	0.350 %
Sorbeth-30	0.400 %
Octyldecanol	0.100 %
Iodopropynyl Butylcarbamate	0.010 %
PEG-4 Laurate	0.090 %
Chamomilla Recutita	0.070 %
Parfum	0.150 %

(continued)

Example 2: aqueous phases	
Phase 2-F	
Tetrasodium EDTA	0.078 %
Citric Acid	0.020 %
Phase 2-G	
Aqua	97.332 %
Phenoxyethanol	0.800 %
Polysorbate-20	0.600 %
PGG-15 Stearyl Ether	0.400 %
Propylene Glycol	0.350 %
Decyl Oleate	0.100 %
Iodopropynyl Butylcarbamate	0.010 %
PEG-4 Laurate	0.090 %
Chamomilla Recutita	0.070 %
Parfum	0.150 %
Tetrasodium EDTA	0.078 %
Citric Acid	0.020 %
Phase 2-H	
Sodium Myreth Sulfate	10.00 %
Lauryl Glucoside	15.00 %
Cocamidopropyl Betaine	10.00 %
Aqua	64.50 %
Parfum	0.50 %
Phase 2-I	
Sodium Laureth Sulfate	20.00 %
Decyl Glucoside	5.00 %
Cocamidopropyl Betaine	8.00 %
Laureth-2	2.50 %
Polysorbate-20	1.00 %
Aqua	63.00 %
Parfum	0.50 %
Phase 2-J	
Sodium Myreth Sulfate	15.00 %
Lauryl Glucoside	10.00 %
Laureth-2	1.50 %
Aqua	73.00 %
Parfum	0.50 %
Phase 2-K	
Emulgade® CM	20.00 %
Polysorbate 20	0.80 %
Coco-Glucoside	2.50 %
Phenoxyethanol	1.00 %
Cetylpyridinium Chloride	0.10 %

(continued)

Example 2: aqueous phases	
Phase 2-K	
Tetrasodium EDTA	0.20 %
Aqua	75.22 %
Citric Acid	0.08 %
Parfum	0.10 %
Phase 2-L	
Emulgade® SE-PF	1.66 %
Cetareth-12	0.94 %
Lamesoft® PO 65	0.25 %
Paraffinum Liquidum	3.00 %
Cetylpyridinium Chloride	0.05 %
Polysorbate-20	1.00 %
Citric Acid	0.03 %
Tetrasodium EDTA	0.20 %
Nipaguard® IPF	0.10 %
Aqua	92.66 %
Parfum	0.11 %
Phase 2-M	
Emulgade® SE-PF	1.627 %
Cetareth-12	0.921 %
Lamesoft® PO 65	0.245 %
Paraffinum Liquidum	2.940 %
Glyceryl Polymethacrylate	2.000 %
Cetylpyridinium Chloride	0.049 %
Polysorbate-20	0.980 %
Citric Acid	0.029 %
Tetrasodium EDTA	0.196 %
Nipaguard® IPF	0.098 %
Aqua	90.807 %
Parfum	0.108 %

Example 3

[0230] Dry hydro-entangled carrier material made of fabric having a surface weight of 50 g/m² was cut into strips. The lipid phase, prepared as set forth in example 1, was applied onto both sides of the fabric with the total amount of 1.0 g in the form of stripes by using the contact process. This comprised running the fabric strips in which the strips against two heated heads having a slitted blade each mounted on one side of the fabric strip. The lipid phase on the strips was allowed to cool so that it solidified and the strips were subsequently sprayed in the conventional manner with the liquid as prepared in example 2. Liquid addition was set at 6 g per wipe. Subsequently the strips were folded and cut.

Claims

1. A product comprising a porous or absorbent sheet whereto an aqueous and a lipid phase have been applied. 5
2. A product according to claim 1 wherein the melting point or melting range of the lipid phase is above or equal to 25 °C. 10
3. A product according to any of claims 2 wherein the melting point or melting range of the lipid phase is in the range of 32 to 40 °C.
4. A product according to any of claims 1 to 3 wherein the lipid phase comprises mono-, di- or triglycerides. 15
5. A product according to claim 4 wherein the lipid phase comprises mono-, di- or triglycerides derived from or present in natural oils. 20
6. A product according to claim 4 wherein the lipid phase comprises fatty acid mono-, di- or triglycerides wherein the fatty acids contain from 12 to 24, preferably from 16 to 20 carbon atoms. 25
7. A product according to claim 4 wherein the lipid phase comprises triglycerides selected from glyceryl stearate, glyceryl oleate, glyceryl laurate, glyceryl myristate, cocoglycerides, or hydrogenated palm oil glycerides, hydrogenated castor oil, or hydrogenated rapeseed oil. 30
8. A product according to any of claims 4 to 7 wherein the lipid phase comprises mono-, di- or triglycerides in an amount of at least 50 %, preferably at least 70 %, more preferably at least 90 %, w/w of the total amount of components making up the lipid phase. 40
9. A product according to any of claims 1 to 3 wherein the lipid phase contains fatty alcohols.
10. A product according to claim 9 wherein the lipid phase contains C₁₂-C₅₀-fatty alcohols, in particular the C₁₂-C₂₄-fatty alcohols. 45
11. A product according to claim 10 wherein the fatty alcohols are selected from myristyl alcohol, 1-pentadecanol, cetyl alcohol, lauryl alcohol, oleyl alcohol, palmityl alcohol, 1-heptadecanol, stearyl alcohol, cetearyl alcohol, 1-nonadecanol, arachidyl alcohol, 1-heneicosanol, behenyl alcohol, brassidyl alcohol, lignoceryl alcohol, ceryl alcohol or myricyl alcohol and C₁₆/C₁₈-Guerbet alcohols. 50
12. A product according to any of claims 9 to 11
- wherein the fatty alcohols are present in the lipid phase, in an amount relative to the total weight amount of the lipid phase, which is in the range of 1 - 40 %, preferably 1 - 30 % (w/w), more preferably of 1 - 20 % (w/w), still more preferably from 1 - 10 % (w/w).
13. A product according to any of claims 1 to 3 wherein the lipid phase contains fatty acids.
14. A product according to claim 13 wherein the fatty acids are C₁₄-C₄₀-fatty acids or in particular are C₁₆-C₃₀-fatty acids.
15. A product according to claim 13 wherein the fatty acids are selected from myristic-, pentadecanoic-, palmitic-, margaric-, stearic-, nonadecanoic-, arachic-, behenic-, lignoceric-, cerotic-, melissic-, erucaic-, elaeostearic-, oleic-, linoleic-, lauric acid and hydroxy-substituted fatty acids.
16. A product according to any of claims 13 to 15 wherein the total amount of the fatty acids present in the lipid phase, relative to the total weight amount of the lipid phase, is in the range of 1 - 30 % (w/w), preferably of 1 - 20 % (w/w), more preferably from 1 - 10 % (w/w).
17. A product according to any of claims 1 to 3 wherein the lipid phase contains one or more of components (a), (b), (c), (d), (e) or (f) as defined hereafter:
 - (a) at least 1 - 50 % (w/w), in particular at least 1 - 10 % of an oily or waxy component
 - (b) 0,1 - 5 % (w/w) of at least one active ingredient
 - (c) 1 - 10 % (w/w) of at least one oil
 - (d) 0.1 - 10 % (w/w) of at least one emulsifier
 - (e) 5 - 90 % (w/w) of further waxy components
 - (f) 0 - 5 % (w/w) water.
18. A product according to claim 17 wherein the lipid phase contains all components (a)-(f).
19. A product according to claim 17 or 18 wherein component (a) is an oily or waxy component selected from C₁₄-C₃₀-dialkyl ethers, C₁₄-C₃₀-dialkyl carbonates, C₄-C₃₄-dicarbonic acids or C₁₂-C₃₀-hydroxyfatty alcohols or mixtures thereof.
20. A product according to any of claims 1 to 3 wherein the lipid phase contains dialkyl(ene) ethers or -carbonates, dicarboxylic acids or hydroxy fatty alcohols, or a combination thereof.
21. A product according to any of claims 1 to 20 wherein the lipid or the aqueous phase contains one

or more active substances.

22. A product according to claim 21 wherein the active substance(s) is or are anti-microbials, e.g. antibacterials and antifungals, anti-inflammatory agents, anti-irritating, anti-itching, anti-perspirant agents. 5

23. A product according to any of claims 1 to 20 wherein the lipid or the aqueous phase contains at least one moisturizer, deodorant, skin caring ingredient, plant extract, vitamin, perfume oil, dye, sunscreen filter, hydrotrope or self-tanning agent. 10

24. A product according to any of claims 1 to 20, wherein the lipid or the aqueous phase contains at least one emulsifier. 15

25. A product according to any of claims 1 to 20, wherein the lipid phase contains at least one superfatting agent, thickener, cationic polymer, anionic polymer, zwitterionic polymer, amphoteric polymer, consistency agent, anti-oxidant. 20

26. A product according to any of claims 1 to 20 wherein the lipid or the aqueous phase contains an insect repellent, a sunscreen filter, a powder or a peeling agent. 25

27. A product according to any of claims 1 to 26 which is a wipe. 30

28. A product according to claim 27 wherein the wipe is a non-woven wipe. 35

29. A method of manufacturing a product as claimed in any of claims 1 to 28 said method comprising contacting the sheet with a lipid phase and with an aqueous phase, either subsequently or simultaneously. 40

30. A method according to claim 29 wherein a lipid phase having a melting point or a melting range of above room temperature is first applied to the surface of the sheet and subsequently the aqueous phase is applied. 45

31. A method according to claim 29 wherein the aqueous phase is applied by spraying, dripping, immersing or running through a bath, and the lipid phase is applied by spraying, contacting, printing or a direct contact process where there is a direct contact between the sheet and an application head having slit nozzles. 50

33. Use of a product as claimed in any of claims 1 to 28 as a combined cleanser and applicator of active substances. 55



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Application Number
EP 01 20 2691

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